



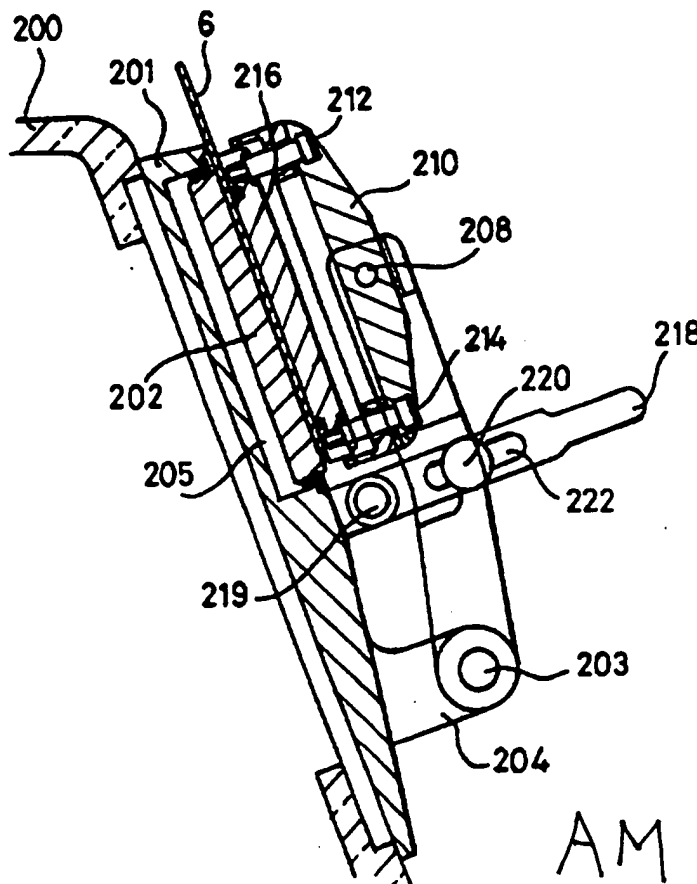
## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>6</sup> : <b>B01L 7/00, G01N 1/31, C12Q 1/68</b>	<b>A1</b>	(11) International Publication Number: <b>WO 96/30124</b> (43) International Publication Date: <b>3 October 1996 (03.10.96)</b>
<p>(21) International Application Number: <b>PCT/GB96/00758</b></p> <p>(22) International Filing Date: <b>27 March 1996 (27.03.96)</b></p> <p>(30) Priority Data: <b>9506312.9</b>      <b>28 March 1995 (28.03.95)</b>      <b>GB</b></p> <p>(71) Applicant (for all designated States except US): <b>MEDICAL RESEARCH COUNCIL [GB/GB]; 20 Park Crescent, London W1N 4AL (GB).</b></p> <p>(72) Inventor; and (75) Inventor/Applicant (for US only): <b>FREEMAN, Thomas, Charles [GB/GB]; 30 Catharine Street, Cambridge CB1 3AW (GB).</b></p> <p>(74) Agent: <b>KEITH W. NASH &amp; CO.; 90-92 Regent Street, Cambridge CB2 1DP (GB).</b></p>	<p>(81) Designated States: <b>AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</b></p> <p><b>Published</b> <i>With international search report.</i></p> <p><b>BEST AVAILABLE COPY</b></p>	

(54) Title: **IMPROVEMENTS IN OR RELATING TO SAMPLE PROCESSING**

## (57) Abstract

Disclosed is a support retaining member (216) for use in processing a sample supported on a support (6), the member (216) being such that when assembled together with a support (6) it forms a support cell, the support cell comprising a substantially sealed chamber, the chamber being provided with a fluid inlet and a fluid outlet for the introduction and removal respectively of fluids used in processing the sample.





**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LI	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Latvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam



**Title:**        Improvements in or Relating to Sample Processing

Field of the Invention

This invention relates to a support cell for processing samples, particularly biological samples, supported on solid supports. In addition the invention relates to automated apparatus for processing samples supported on solid supports, which automated apparatus may be used with the support cell of the invention. The invention also relates to a method of processing a sample supported on a solid support.

Background of the Invention

It is common practice, in certain technical fields, for samples of interest to be subjected to a particular process to be supported on solid supports. For example, protein or nucleic acid samples may be supported on filters or membranes: typically such filters or membranes are made of nylon or nitrocellulose. Samples, especially biological samples (e.g. cell suspensions or tissue sections) may also be supported on slides. Typically the slides are of transparent glass.

Samples supported on slides may be processed by a number of various techniques known to those skilled in the art, which include: standard staining techniques (e.g. Gram staining of micro-organisms, haemotoxylin/eosin staining of tissue sections); immuno- and enzyme cytochemistry (e.g. peroxidase/anti-peroxidase staining); ligand binding studies; in situ hybridisation of labelled DNA or RNA probes; and in situ amplification of nucleic acids present in a sample by means of polymerase chain reaction (PCR).

Typically, such processing is at present performed manually by laboratory personnel. This renders the processing slow and subject to individual variation (from one sample to the next, and/or from one day to the next), and is tedious for the personnel concerned. Some attempts have been made to provide apparatus to facilitate the immuno-cytochemistry



process. Thus, for example, Shandon Scientific Limited manufacture the "Sequenza"<sup>TM</sup> and "Cadenza"<sup>R</sup> immuno-staining apparatus. Use of the "Sequenza"<sup>TM</sup> apparatus involves attaching a coverplate to the slide to be processed, thereby forming a narrow channel between the slide and coverplate. The coverplate comprises a funnel-like open-ended portion which, when attached to a slide, forms an open-ended chamber, into which staining reagents are manually introduced. The reagent then flows, under the influence of gravity, into the channel between the slide and the attached coverplate. When a new reagent is introduced into the well it displaces the previous reagent into a waste collection tray at the bottom of the apparatus.

The "Cadenza"<sup>R</sup> apparatus utilises the same coverplate system but additionally provides programmable automated control means for applying reagents to the open-ended well of the assembled slide/coverplate.

In relation to processing of samples supported on membranes, such samples are commonly preparations of proteins (termed Western blots) or DNA (Southern blots) or RNA (Northern blots). The molecular species of interest is visualised on the supporting membrane using specific antibodies and an appropriate detection systems (in the case of Western blots) or by hybridisation with a labelled probe (in the case of Southern and Northern blots). These processing techniques currently require considerable manual processing of the membranes.

The prior art apparatus for processing samples on solid supports is therefore very simple and limited in the functions which it can perform. The present invention aims to provide apparatus capable of more general application.

#### Summary of the Invention

In a first aspect the invention provides a support retaining member for use in processing a sample supported on a support, the member being such that when assembled together with a support it forms a support cell, the support cell comprising a substantially sealed chamber, the chamber being provided with a fluid inlet and a fluid outlet for the



introduction and removal respectively of fluids used in processing the sample.

In a particular embodiment, part of the internal surface of the substantially sealed chamber is defined by the sample bearing surface of the support, which arrangement has the advantage of minimising the number of components involved.

Conveniently the support is a slide, or a membrane of the type known to those skilled in the art.

Where the support is a slide, the support retaining member will typically be of such dimensions that it may be used with slides of conventional size, (i.e. slides which typically are about 25.4mm by 76.2mm). Where the support is a membrane, the dimensions of the retaining member will similarly be of such dimensions that it may be used with membranes of conventional size (typically 80mm by 120mm), although membranes are rather more variable in size than slides.

In some embodiments the support retaining member comprises two opposed portions, the support being positioned between the opposed portions of the support retaining member when the support cell is assembled. Preferably at least that portion of the support retaining member which comes into contact with the fluids used to process the sample on the support is made of an inert material, such as a synthetic moulded plastics material. Perspex is found particularly suitable. The remainder of the support retaining member may generally be of the same or other material, e.g. glass, perspex or metal (such as duraluminium).

The components of the support cell are typically held together by clamping means. In certain embodiments the clamping means is capable of clamping together the portions of a plurality of support cells. Typically, from one to around twelve or sixteen support cells may be clamped simultaneously by a single clamping means. The support cells may preferably be arranged in the clamping means in a substantially horizontal or substantially vertical manner, although any position intermediate between these two positions may be possible. The support cells may, for example, be arranged side-by-side, or one in front



of the other.

As an alternative, or in addition, to clamping means the support cell may be provided with biasing means, which biasing means tends to urge together the components of the support cell. Conveniently the biasing means will comprise one or more sprung biasing members. In a particular embodiment, the support retaining member is attached to clamping means by spring-loaded mounting pins, such that formation of the support cell places the springs of the spring-loaded mounting pins under compression, which springs therefore urge together the components of the support cell.

In preferred embodiments, the force applied to the support cell by the clamping means and/or the biasing means helps to ensure a fluid-tight seal between the support and the support retaining member.

It is generally preferred that the support cell additionally comprises sealing means to assist in the formation of the substantially sealed chamber. Conveniently the sealing means forms a fluid-tight seal, in the first instance with the sample bearing surface of the support and, in the second instance, with the support retaining member. Alternatively, the sealing means may effect a seal directly between opposed portions of the support retaining member, where the support retaining member is of two-part construction. The sealing means may be an integral part of the support retaining member, or may be provided as a separate component of the support cell. The sealing means typically comprises a gasket, which may be made of silicon rubber or other suitable material. In one embodiment the sealing means comprises an O-ring gasket, the shape of which is generally that of a frame-like surround seated in a groove in one portion of the support retaining member. In an alternative embodiment the sealing means comprises a flattened frame-like surround gasket (about 100 to 150 $\mu$ m thick). Either type of gasket may be discarded after a single use (if, for example, contaminated with a radioactive probe) or may be re-used if desired. The flattened gasket embodiment is particularly suitable as a disposable gasket, to be discarded after a single use. It will be apparent that the thickness of the gasket (which can be readily altered by exchanging gaskets) may, in part, determine the volume of the substantially sealed chamber.



Conveniently the fluid inlet into, and the fluid outlet out of, the substantially sealed chamber are formed in the support retaining member. Where the support is a slide, the fluid inlet and fluid outlet are preferably provided in one portion where the support retaining member is of the two-part construction defined above. Where the support is a membrane, however, it is preferred that the fluid inlet and fluid outlet are provided in opposed portions of the support retaining member, where the support retaining member is of two-part construction. This encourages correct flow of processing fluids through the membrane.

The fluid inlet allows the introduction into the substantially sealed chamber of fluids needed to process the sample on the support. Typically such fluids will be buffers, solvents (e.g. ethanol/methanol, xylene), reagents (e.g. antibody- or probe-containing solutions) or stains. The fluid outlet allows for the processing fluids to be removed from the sample (e.g. for a washing step, or to allow the addition of a further reagent). Preferably, when the supports are being processed, their orientation is such that the fluid inlet is in the bottom portion of the substantially sealed chamber, and the fluid outlet is in the top portion of the substantially sealed chamber.

Typically, where the sample is supported on a slide, the substantially sealed chamber will have a volume of between  $50\mu\text{l}$  and  $300\mu\text{l}$ , preferably between  $100\text{--}150\mu\text{l}$ . This small volume allows for economical use of reagents and (where temperature regulation is involved) a rapid thermal response time. Where the sample is supported on a membrane, the chamber will generally be larger (up to 2-3mls).

In a particular embodiment, the support cell is adapted so as to be suitable for use in performing PCR on samples supported on a support (typically a slide), which hitherto has proved extremely difficult: performing PCR involves heating small volumes of liquids to comparatively high temperatures, which conventionally necessitates performing the PCR in a sealed reaction chamber to prevent undue evaporation loss of the reactants. However, the sealed chamber must be openable to allow the addition of further reagents. Such an openable chamber has hitherto been very difficult to create in the context of a standard slide. At present, the sealed chamber is typically created by placing a glass coverslip over



a volume of PCR mix and sealing around the coverslip with a substance such as mineral oil or nail varnish. Re-opening the chamber is difficult and contamination of the chamber contents sometimes occurs. Also, the presence of the slide has hindered the ability rapidly to alter the temperature of the reagents, which ability is desirable when performing PCR.

In a particular embodiment envisaged for PCR (or other processes where temperature regulation is required), the support cell is equipped with temperature control means to allow for rapid heating and cooling of the sample and PCR mix (i.e. thermal cycling). Typically the support cell will be provided with an electrical heating element or a Peltier device. The support cell may also be adapted (e.g. by provision of cooling fins) to provide for improved air cooling. Temperature control in the range 4°-100°C is found sufficient for most applications.

In a further aspect, the invention provides automated apparatus for processing a sample, especially a biological sample, supported on a support, the apparatus comprising: support holding means for holding one or more supports, the sample on the or each support being present within a respective substantially sealed chamber; fluid delivery means for delivering processing fluid to the or each chamber; waste fluid collecting means for removing fluid from the or each chamber; and computer control means.

Preferably the apparatus is used in conjunction with the support cell defined above. Conveniently the holding means comprises clamping means suitable for clamping together the components of the support cell, as defined previously. Conveniently the holding means can accommodate from ten to twenty-four support cells.

A number of arrangements for appropriate fluid delivery means can be envisaged. In a preferred embodiment a number of reservoirs (typically, 10) of processing fluids, (e.g. buffers, stains, etc.) are provided, each reservoir being attached to pump means. Preferred pump means are syringe pumps, such as those manufactured by Hook and Tucker, (Croydon, Surrey, UK), or Kloen having a stroke volume of between 1 and 10ml. One such pump may be provided for each processing fluid reservoir, or a single pump may be provided to pump fluid from each a plurality of reservoirs, by means of a multi-



port valve configuration.

Conveniently each syringe pump is in turn attached to a central manifold (such as a universal connector). Preferably the central manifold feeds into a selective multi-outlet valve such that, if desired, where a plurality of samples are being processed simultaneously, each sample may be treated with a different processing fluid or combination of processing fluids, such that samples can be individually processed. A suitable selective multi-outlet valve is a rotary valve, such as the 10 outlet rotary valve supplied by Omnifit (Cambridge, UK). Thus each outlet from the multi-outlet valve may be connected to a separate support cell. One or more filters (especially "on-line" filters) may be incorporated if desired. Typically a filter will be positioned between each reservoir and its associated syringe pump.

Each syringe pump may be actuated individually by the computer control means, or two or more pumps may be actuated simultaneously to provide a mixture of two or more processing fluids. Controlling the rate of operation of each pump will thus control the composition of the resulting mixture of processing fluids.

In an alternative embodiment, the fluid delivery means comprises two or more piston/HPLC-type pumps, each pump being supplied, via a multi-inlet valve, by a plurality of processing fluid reservoirs. Suitable pumps are available, for example, from Anachem (Luton, Beds, UK). Conveniently the multi-inlet valve will be a rotary valve. Each pump will feed into a rotary mixer, of the type well known to those skilled in the art, thus allowing variable composition mixtures of processing fluids to be produced, if desired.

Preferably, the processing fluid or mixture of processing fluids is then passed through an in-line filter and then conveniently passes through a selective multi-valve outlet (such as a rotary valve) before being fed into the support cells. Conveniently, each support cell inlet will be provided with a respective valve.

As an alternative to the generally "parallel" supply of processing fluids defined above, the processing fluids may be supplied in "series" such that, for example, fluid is passed from



one substantially sealed chamber to another. This embodiment would be particularly suitable, for example, when applied to simple staining of samples and has the advantage that the amount of reagent required is minimised.

In general relation to the fluid delivery means, it is a further preferred feature that a multi-inlet valve be located close to the fluid inlet provided in the or each substantially sealed chamber. Typically the valve will be a three-way valve with two inlets, and one outlet leading to the substantially sealed chamber. One of the valve inlets is fed, indirectly, by the reservoirs of processing fluid. The second inlet is conveniently fed by a local reservoir which, typically, will be a syringe, pipette or micro-pipette (generally 100-5000 $\mu$ l volume). This local reservoir may be controlled by the computer control means or may be manually controlled. The local reservoir will typically be used where a reagent is scarce (e.g. an antibody) or where the reagent is radioactive (e.g. a radio-labelled probe or ligand). The provision of such a local reservoir minimises the amount of reagent required, simplifies radioactive decontamination procedures, and provides extra flexibility in that each support cell may be processed individually, if required.

Preferably processing fluid will enter the or each respective substantially sealed chamber at the bottom, travel upwards and exit from the chamber via the fluid outlet at the top. The or each fluid outlet will conveniently empty into a common collecting duct, which duct drains into a collecting vessel. The vessel is desirably removable from the apparatus to allow for periodic emptying or decontamination.

In a preferred embodiment, the apparatus is also equipped with temperature control means, as many procedures require some or all of the steps to take place at a specific temperature (e.g. binding of antibodies, hybridisation of probes and primers or labelled ligands). These temperature dependent steps are generally performed between 4°C and 95°C, for between 30 minutes and 24 hours. The temperature control means may be incorporated into the support cells (e.g. in the form of heating elements). Alternatively or additionally the support holding means may comprise, or be formed with, or be situated within, temperature control means. In a particular embodiment the support holding means is mounted on a block provided with a Peltier device temperature control means.



Additionally, a fan may be provided to assist cooling of the apparatus.

The computer control means will generally comprise a personal computer (PC) or, more preferably, the computer control means will be integrated within the automated apparatus. Ideally, the computer control means should control two or more (preferably all) of the following parameters: the selection of which pump or pumps to actuate; the absolute volume and the rate of flow of processing fluid passing through the actuated pump(s); the selection of which support cell to feed with processing fluid; the temperature of the supported samples within the apparatus; and the timing of the various events.

It is envisaged that apparatus in accordance with the invention will be suitable for use, inter alia, in standard staining techniques, immuno- and enzyme cytochemistry (e.g. peroxidase/antiperoxidase staining); ligand binding studies; in situ hybridisation of labelled DNA or RNA probes; or even in situ PCR and in processing of Southern, Northern or Western blots. Preferably the computer control means has a programmable memory such that the steps needed to perform a certain procedure (e.g. haemotxylin/eosin staining) can be entered into the computer memory and readily re-called without the need to re-program the computer.

Conveniently the apparatus may be substantially modular such that, should large numbers of supported samples require to be processed, further reservoirs, pumps, support cells etc., can be readily added to the existing equipment. In such an embodiment the apparatus is preferably capable of accepting a modular array of support cells, whether the samples are supported on slides or membranes.

The invention further relates to manufacture of and use of the support cell and/or the apparatus of the invention in processing a sample on a support. such that the invention provides: a method of processing a sample on a support using a support cell and/or the automated apparatus defined above; a method of making a support cell and a method of making automated apparatus in accordance with the invention.

The invention will now be further described by way of illustration and with reference to



the accompanying drawings in which:

Figure 1a is a plan view of a portion of a support retaining member (where the sample support is a glass slide), Figures 1b and 1c being side and end elevations respectively of the portion illustrated in Figure 1a; Figure 1d being a plan view of the portion of a support retaining member opposed to that shown in Figures 1a-1c, Figure 1e being an end elevation of the portion shown in Figure 1d, Figures 1f and 1g being respective sections along the lines XX and YY shown in Figure 1d;

Figure 2 is a plan view of a portion of a support retaining member, being a different embodiment to that shown in Figure 1a;

Figures 3a and 3b are side and plan elevations respectively of an array of support cells in accordance with the invention clamped within clamping means, which clamping means also forms part of support holding means in apparatus in accordance with the invention;

Figure 4 shows a perspective view of apparatus in accordance with the invention and Figure 5 shows a perspective view of a further embodiment of apparatus in accordance with the invention;

Figure 6 is a schematic representation of part of the fluid delivery means for use in one embodiment of the apparatus of the invention;

Figures 7a and 7b are longitudinal sections through a support cell in accordance with the invention, together with a schematic representation of part of the associated fluid delivery means, the plane of the section in Figure 7b being at 90° to the plane of the section in Figure 7a;

Figure 8a is a flow diagram illustrating schematically the steps performed in haemotoxylin/eosin staining of samples and Figure 8b shows how the same process could be performed using apparatus in accordance with the invention;



Figures 9a-9c are flow diagrams illustrating schematically how apparatus in accordance with the invention could be used to perform immunocytochemistry processing, in situ hybridisation and in situ PCR respectively;

Figures 10a-10b are enlarged, exploded sectional and perspective views respectively of a support cell wherein the sample is immobilised on a membrane;

Figure 11a is side elevation (to a different scale) of apparatus in accordance with the invention, and Figure 11b is a sectional view of the same, along the dotted line Y-Y in Figure 11a;

Figure 12 shows, on a different scale, a sectional view of apparatus in accordance with the invention, similar to that shown in Figure 11, adapted for performing processing of samples requiring careful temperature regulation (e.g. in situ PCR);

Figures 13a and 13b show in cross-section apparatus in accordance with the invention wherein sample supports are positioned substantially horizontally;

Figures 14a-14c show, in cross-section, sample retaining members suitable for use with particular embodiments of apparatus according to the invention, Figure 14d shows a plan view of a further sample retaining member, and Figure 14e is a cross-section of the member shown in Figure 14d;

Figures 15a and 15b are side elevations (partly in sectional view) of apparatus in accordance with the invention, in an open, disassembled position (15a) and in a closed, assembled position (15b);

Figure 16 is a front elevation of an array of six support cells of the type shown in Figure 15, in their assembled position; and

Figure 17 is a schematic representation of fluid delivery means for use with apparatus in accordance with the invention.



### Detailed Description of Embodiments of the Invention

In the embodiment shown in Figures 1a-g, a support cell for use in processing a sample on a slide comprises a support retaining member 2 and sealing means 4, the support retaining member 2 being such that when assembled together with a support 6 and sealing means 4 it forms a support cell, the support cell comprising a substantially sealed chamber 8 the internal surface of which is defined in part of the sample bearing surface of the support 6, the chamber 8 being provided with a fluid inlet 10 and a fluid outlet 12 for the introduction and removal respectively of fluids used in processing the sample. The support 6 is a conventional glass slide.

The support retaining member 2 comprises two opposed portions, 2a and 2b, shown in plan view in Figures 1a and 1d respectively. The support 6 is positioned between the opposed portions 2a and 2b when the support cell is assembled.

Referring to Figure 1a, one portion 2a of the support retaining member 2 is a generally rectangular block of perspex, about 37mm wide, 7.5mm thick and 60mm long. In addition, the block is formed with a central raised region 7, 29mm wide, 2.5mm high and about 55mm long. This central raised region is clearly shown in Figures 1b and 1c.

The sealing means 4 is a flattened silicon rubber gasket in the form of a frame-like surround, about 4mm wide. The gasket 4 is of sufficient dimensions to extend around the fluid inlet 10 and the fluid outlet 12.

The fluid inlet 10 takes the form of a cylindrical bore, with a diameter of about 1mm. The fluid inlet 10 enters the portion 2a of the support retaining member at one side (as shown in Figure 1b). The entrance to the fluid inlet 10 is formed with a cylindrical counter-bore about 10mm deep and about 5.6mm in diameter. The counter-bore facilitates the attachment of fluid delivery means. The cylindrical bore of the fluid inlet extends into the portion 2a about 20mm and then passes through 90° to form an opening at one end of the central raised region of the portion 2a. The cylindrical bore of the fluid inlet 10 at this



point widens into a long slot 14, extending across most of the width of the central raised region.

The other end of the central raised region has a larger slot 16 which narrows into the opening of the fluid outlet 12. The fluid outlet 12 is a cylindrical bore with a diameter of 2mm (about twice the diameter of the fluid inlet 10), which passes through 90° to exit the portion 2a of the support retaining member at the same side as that on which the fluid inlet 10 enters the portion 2a (see Figure 1b). The fluid outlet 12 is counter-bored in a similar manner to the fluid inlet 10 in order to facilitate attachment of waste fluid removal means.

As shown in Figure 1d, the portion 2b of the support retaining member 2 is a generally rectangular block of duraluminium with overall dimensions similar to the portion 2a of the support retaining member 2. The portion 2b has stepped sides 19 and a stepped end region 18 at one end. Figure 1e shows an end elevation of the portion 2b as seen from the end with the stepped end region 18 (the stepped sides are omitted for clarity). Figures 1f and 1g are sectional views along the lines XX and YY (in Figure 1b) respectively. Figures 1a - 1g are all drawn to the same scale.

In use, a support 6 (glass slide) bearing a sample to be processed is placed in the portion 2b of the support retaining member 2, the stepped sides 19 and stepped end region 18 serving to guide and retain the slide in the correct position. The support 6 is positioned such that the surface of the support 6 bearing the sample is furthestmost from the portion 2b of the support retaining member 2.

The sealing means 4 (gasket) is then placed around the uppermost surface of the support 6, surrounding the sample to be processed. The positioning of the sealing means 4 is assisted by the stepped edges 19 and stepped end region 18 of the portion 2b of the support retaining member. The portion 2a of the support retaining member is then positioned on top of the sealing means 4, so as to make a substantially sealed chamber 8 largely defined by the surface of the support 6 bearing the sample and the opposed surface of the raised central region 7 of the portion 2a of the support retaining member 2. Again,



positioning of the portion 2a is assisted by the stepped edges 19 and stepped end region 18 of the portion 2b. The apertures of the fluid inlet 10 and the fluid outlet 12 are within the substantially sealed chamber 8, the edges of which are defined by the sealing means 4.

The support cell assembly is then placed in a clamping means, ready for processing.

Figure 2 shows a portion 2c of a support retaining member. The portion 2c is a generally rectangular perspex block substantially similar to the portion 2a shown in Figure 1a. However, in the embodiment shown in Figure 2, the sealing means 4 is a silicon rubber O-ring, which is seated in an accommodating groove cut in the raised central region of the portion 2c. The accommodating groove is about 1.5mm wide and 0.88mm deep. The mean corner radius of the turns in the groove at the four corners is about 3mm.

Figure 3a is a side elevation of clamping means 20 securing an array of eight support cells of the type shown in Figures 1 or 2. Figure 3b shows a plan view of the same array. The clamping means 20 is generally rectangular and comprises a fixed end block 22, 24 at opposed ends, between which is provided a shallow channel (a few millimetres deep), of a width sufficient to accommodate a support cell array. The channel is slightly off-set from the medial axis of the clamping means. Fixed block 24 is formed with a screw-thread aperture, which aperture is in screw-threaded engagement with clamping screw 26. The outer end of clamping screw 26 is knurled to facilitate grip by an operator's fingers (but in an alternative embodiment a rod may be inserted through an aperture provided in the outer end of the clamping screw, perpendicular to the long axis thereof, which rod can be used to increase the turning moment applied to the screw). The inner end of clamping screw 26 is attached to a clamping face 28. Accordingly, turning clamping screw 26 moves the clamping face 28 towards or away from the fixed end block 22, such that between one and eight support cells may be secured in the clamping means 20 with a variable clamping force, which clamping force, in addition to securing the support cells also serves to ensure a good seal between the sealing means and the support/support retaining member.



The clamping means 20 is made of metal and has a perspex cover (not shown), which serves to prevent excessive heat loss of samples and is a safety feature when radioactive substances are used in the processing of the sample.

The clamping means 20 is about 225mm long, 95mm wide and 50mm thick. The shallow channel is about 38mm wide. However, where the samples on the supports are to be subjected to in situ PCR the overall dimensions of the clamping means will be reduced, thus decreasing its thermal capacity, allowing for quicker thermal cycling.

Typically the clamping means will be compatible with the automated apparatus of the invention and generally the clamping means will form at least part of the support holding means in the automated apparatus.

Figure 4 is a perspective view (to a different scale) of one embodiment of automated apparatus for processing a sample on a support in accordance with the invention. In Figure 4, an array of ten support cells is held in a support holding means 30, which holding means comprises the clamping means 20 of Figure 3. The sample on each support is present within an individual substantially sealed chamber, such as that indicated in Figure 1a. The apparatus also comprises fluid delivery means (indicated generally at 32 in Figure 4) for delivery processing fluid to each chamber, waste fluid collecting means 34 for removing fluid from each chamber, and computer control means 36.

More specifically, the fluid delivery means 32 comprises six reservoirs 38, each reservoir 38 containing about 1 litre of a processing fluid. Fluid is fed from each reservoir 38 through narrow synthetic plastics tubing 39 to an associated syringe pump 40. The syringe pumps are those supplied by Hook & Tucker (Croydon, Surrey, UK) and have a stroke volume of 1-10mls. One syringe pump 40 is provided for each reservoir 38. Each syringe pump 40 feeds via a short length of tubing 41 into a universal connector 42, which is in turn connected via tubing 43 to an Omnifit rotary valve 44, provided with ten outlets. Each outlet from the rotary valve 44 is connected via tubing 45 to a fluid inlet in one of the support cells held in the holding means 30. Each of the fluid outlets is connected to a common waste fluid collecting means 34 which drains into a glass beaker 46.



The apparatus further comprises a perspex cover 48 located over the array of support cells. The support holding means 30 is located on a temperature control means 50, which takes the form of a fan-assisted Peltier device (fan omitted for clarity). A variable temperature regulation control 52 is provided. The computer control means is a PC which controls: the selection of which syringe pumps 40 to actuate; the absolute volume and the rate of flow of processing fluid; the selection of which support cell to feed with processing fluid; the temperature of the supported samples within the apparatus; and the timing of the various events.

An alternative embodiment of the apparatus is shown (to a different scale) in Figure 5. The embodiment shown in Figure 5 comprises two support holding means 60, each of which is capable of supporting up to sixteen supports 6. The holding means 60 are mounted on a programmable thermal cycler 62. One of the holding means is occupied by eleven slides 6 and is covered with a perspex lid 66. Five reservoirs 68 of processing fluid are arranged on a shelf. Two reservoirs 68 are connected in a synthetic plastics tubing 69a to rotary valve 70a whilst the other three are connected via tubing 69b to rotary valve 70b. Rotary valve 70a is in turn connected via tubing 71a to HPLC pump 72a, whilst rotary valve 70b is connected through tubing 71b to HPLC pump 72b. The HPLC pumps 72a, 72b each feed through tubing 73a,b into mixer unit 74, thus allowing variable composition mixtures of processing fluids to be produced, if desired. The processing fluid(s) passes through tubing 75 and then enters each chamber in the array of support cells and waste fluid exhausted therefrom is collected by a common waste duct 76, which empties into a glass collecting beaker 78.

The arrangement of the fluid delivery apparatus in this embodiment is shown schematically in Figure 6. In Figure 6, a filter unit 80 is included downstream of the mixer unit 74.

In the embodiment of a support cell shown in Figure 7a, a support retaining member 82 accommodates a support 6 (glass slide) and an electrical heating element 84. A substantially sealed chamber 86 is defined by the surface of the support 6 upon which the sample is located and the opposed surface of the support retaining member 82, the seal being effected by a silicon rubber gasket (not shown). A fluid inlet is provided at the



bottom of the chamber 86, which fluid inlet is attached to a three-way valve 90.

In Figure 7b, the main feed from the reservoirs of processing fluid is indicated at 92 and is attached to one of the ports of the three-way valve 90. In addition, a syringe 94 forms a local reservoir which is connected to a second of the ports of the three-way valve 90. The local reservoir 94 will typically be utilised where the processing of the sample involves the use of scarce and/or radioactive reagents. The fluid inlet 88 splits into a number of channels 89, such that processing fluids will enter the substantially sealed chamber 86 along a broad front and advance upwards evenly before exiting the chamber 86 via outlet 96, thence into the common waste collecting duct (not shown).

Figure 8a shows the flow diagram for performing haematoxylin/eosin staining of biological samples.

Figure 8b shows how the process could be performed using automated apparatus in accordance with the invention, the fluid delivery means comprising two HPLC-type pumps.

Similarly, Figures 9a-9c are flow diagrams showing how apparatus in accordance with the invention, utilising an HPLC-type two pump system, could be employed to perform an immunocytochemical process (9a), in situ hybridisation (9b) or in situ PCR (9c). Tables I and II indicate, respectively, sample protocols for performing haematoxylin/eosin staining and in situ hybridisation of a paraffin-treated thin section, using a six-syringe pump system.

In the embodiment shown in Figures 10a and 10b a support cell comprises two opposed portions 96, 98 of a support retaining member. The portion 96 is equipped with a fluid inlet 100 and a recessed area 102. The portion 98 is formed with a fluid outlet 104 and a raised section 106. The dimensions of the recessed section 102 of the portion 96 are such as to accept the raised section 106 of the portion 98, with a membrane sample support 108 positioned therebetween.



The raised section 106 is provided with a peripheral groove, which groove accommodates an O ring silicon rubber gasket 110.

When the support cell is assembled, a substantially sealed chamber is defined by the O ring gasket 110 and the opposed portions 96, 98 of the support retaining member.

A further embodiment is illustrated in Figures 11a-11b.

In Figure 11a a substantially vertical array of 16 support cells 112 is provided, eight support cells 112 being disposed on either side of a central support holding member 114. The sample supports 6 are omitted (for clarity) from the support cells 112 except from that on the left hand end.

The arrangement is more clearly shown in the sectional view of Figure 11b. The sample supports 6 (glass slides) are held on a shelf on either side of the support holding member 114. Each support cell comprises a sample support 6 and a perspex support retaining member 116. The support retaining members 116 are seated on a shelf either side of the support holding member 114, which comprises duraluminium, a good thermal conductor. Each support retaining member 116 is similar to the portion 2a of the support retaining member shown in Figures 1a-1c. Each section of the support holding member 114 which holds a sample support 6 is similar to the portion 2b of the support retaining member shown in Figures 1d-1g. A substantially sealed chamber is defined by the sample bearing surface of the support 6, the opposed surface of the support retaining member 116 and by an O ring gasket sealing means 118. Each substantially sealed chamber is provided with a fluid inlet 120 near the bottom of the support cell, and with a fluid outlet 122 near the top of the support cell. The fluid inlet 120 and outlet 122 are thus provided in a single support retaining member 116. The support cells 112 are clamped in place on the support holding member 114 by clamping means. The clamping means comprises, for each support cell 112, a screw threaded clamping member 124, which is in screw threaded engagement with an engagement bar 126 which runs the length of the apparatus and is supported at each end. The outer end of the clamping member 124 is provided with a knurled knob to facilitate manual operation. The opposite end of the clamping member



124 is provided with, or contacts, a clamping plate 128. Clamping plate 128 serves to spread the clamping force of the clamping member 124 over a greater area of the support retaining member 116.

Figure 12 is a sectional view of similar apparatus in an embodiment adapted to perform processing steps requiring careful temperature regulation, in particular, thermal cycling such as is necessary to perform in situ PCR. The arrangement is similar to that shown in Figure 11b with each support cell comprising a sample support 6 and a perspex support retaining member 116. A substantially sealed chamber is defined by the sample bearing surface of the support 6, the opposed surface of the support retaining member 116 and by an O ring gasket sealing means 118. As previously, the support cells are clamped on either side of the support holding member 114 by clamping members 124 provided with clamping plates 128, the clamping members 124 being in screw-threaded engagement with an engagement bar 126.

The fluid inlet 120 is equipped with a Luer fitting 130, to which is attached a three-way valve 132. The valve 132 allows introduction of processing fluid from remote fluid delivery means via duct 134 which passes through the base of the holding member 114, or introduction of fluid from a local reservoir via short duct 136, which is adapted to receive a micropipette tip. In addition, duct 136 may be sealed at its outer end by a cap (omitted for clarity) or by covering with mineral oil.

The fluid outlet 122 is similarly provided with a luer fitting 138, allowing for ready attachment of waste fluid collecting tubing 140, which passes through the centre of the holding member 114, for convenience.

The support holding member 114 is formed with a plurality of integral heating elements 142 which come into direct contact with the sample supports 6. The heating elements 142 are metallic and electrically powered. The support holding member 114 is also provided with cooling means, which comprises a series of conducting vessels 144 which pass in close proximity to the sample supports 6. The conducting vessels 144 are adapted to receive a rapid flow of a fluid coolant (such as water), which is forced through the



conducting vessels by a pump. The cooling system may be "closed" (with continuous recycling of coolant) or "open" (in which coolant passes out of the system after completing one circuit, and is replenished with fresh coolant).

Figures 13a and 13b illustrate an embodiment of apparatus in accordance with the invention, wherein the support cells are positioned substantially horizontally. As with the embodiments illustrated in Figures 11 and 12, an array of support cells is positioned on a support holding member 114, each support cell comprising a substantially sealed chamber defined by the sample bearing surface of a support 6, the opposed surface of a support retaining member 116 and by an O ring gasket sealing means 118. Positioning of the support 6 and the support retaining member 116 on the support holding member 114 is assisted by a stepped edge 115 provided on the holding member 114.

The support cells are clamped in position on the holding member 114 by clamping means comprising a clamping member 124 formed with a clamping plate 128, the clamping member 124 being in screw threaded engagement with an engaging member 126.

The holding member 114 is provided with a temperature regulation Peltier device 146 which is in direct contact with the sample support 6 and provides for the heating and cooling thereof. In the embodiment shown in Figure 13a, the fluid outlet 122 is equipped with a luer fitting for the convenient attachment of a waste fluid collecting tubing 148, whilst the processing fluid inlet 120 is provided with a luer fitting to which is attached a three way valve 132. The valve 132 allows for the introduction of processing fluid from a remote fluid delivery means (via duct 150) or from a local reservoir, by the insertion of a micropipette tip into duct 152. The duct 152 may be sealed by a synthetic plastics cap (not shown) or by the introduction of mineral oil. The arrangement shown in Figure 13b differs in that the fluid inlet 120 is adapted to receive processing fluid directly from a micropipette tip.

It will be apparent that the holding member 114 in this embodiment could readily be altered by substituting the Peltier device 146 with separate heating and cooling means, as provided in the embodiment shown in Figure 12.



Figure 14 shows a number of various configurations that might be used for the slide retaining member 116 for use with any of the apparatus embodiments shown in Figures 11-13. Figure 14a shows the basic arrangement, with a fluid inlet 120 provided near the bottom of the member 116 (is positioned vertically) and a fluid outlet 122 provided near the top. The member 116 has a stepped profile on its inner face, to assist in position relative to a sample support and/or a support holding member. The member 116 is also provided with an O ring gasket sealing means 118, seated in a peripheral groove. In the embodiment shown in 14b, the fluid inlet 120 is formed with a luer fitting 130; to facilitate the attachment of associated fluid delivery means such as tubing or valves. In the embodiment shown in Figure 14c, both the fluid inlet 120 and the fluid outlet 122 are formed with luer fittings.

The arrangement shown in 14d is slightly more complex in that, when assembled together with an appropriate sample support (e.g. glass slide), the support retaining member is capable of simultaneously forming three substantially sealed chambers 8a, 8b and 8c, each of which is provided with a respective fluid inlet 120a,b,c and a respective fluid outlet 122a,b,c, the sides of the substantially sealed chamber being sealed by respective O rings 118a,b,c.

Figures 15a, b illustrate a preferred embodiment of apparatus in accordance with the invention, as shown in side elevation (partly in sectional view). Figure 15a shows the apparatus in an open position, whilst Figure 15b shows the apparatus in a closed, operational position.

Referring to Figure 15a, the apparatus comprises a perspex housing 200, accommodating a detachable aluminium fitting 201. Four such fittings 201 are provided on the apparatus, although only one is shown, the rest being omitted for clarity. The fitting 201 detaches from the housing 200 to facilitate cleaning, maintenance and the like. Each fitting 201 is formed with six metallic support holding members 202. Each support holding member 202 is retained in the fitting 201 by a silicone rubber gasket in such a way that an air-filled space 205 is formed between the support holding members 202 and the fitting 201. Each support holding member 202 is equipped with electrical heating means. The



insulation afforded by the air-filled space 205 allows for rapid temperature regulation of the support holding members 202 by their respective electrical heating means, and thus rapid temperature regulation of supports held on the support holding members 202. Each support holding means 202 is also provided with temperature sensing means (not shown).

Twelve projections 204 are provided on the fitting 201, one pair of projections 204 being located beneath each support holding member 202. A pair of angled arms 206 is pivotally mounted at one end region on a pivot 203 provided in each pair of projections 204. At the opposed end region of the angled arms 206 is provided a pivotal attachment 208, upon which is mounted a pivoting head-piece 210. The head-piece 210 is provided with an upper and lower spring-loaded mounting pin 212, 214, which pins 212, 214 are engaged with a support retaining member 216.

There are thus six support retaining members 216 associated with each of the four fittings 201 provided on the apparatus. Each support retaining member 216 is of substantially similar construction to the portion labelled 2a of the support retaining member shown in Figures 1a-1c. The support retaining member 216 is formed in perspex with a fluid inlet and a fluid outlet (not shown in Figure 15, but labelled as 228 and 230 respectively in Figure 16).

The arrangement is such that the pair of angled arms 206 may be moved about pivot 203 in a vertical plane from an open position, as shown in Figure 15a, to a closed, operational position as shown in Figure 15b, and vice versa. Movement of the angled arms 206 upwards towards the housing 200, brings the support retaining member 216 into close proximity, in the closed position, with the support holding member 202. Movement of the angled arms 206 may be effected manually, by the operator of the apparatus.

The apparatus may be retained in the closed position by actuation of a clamping means, which comprises a metallic, generally cylindrical, clamping lever 218 provided between each pair of angled arms 206 and pivotally mounted, at one end region, to a pivot 219 provided on the housing 200 above the projection 204. Within the clamping lever 218 is a transverse member 220. The transverse member 220 protrudes on each side beyond the



surface of the clamping lever 218, and has limited degree of travel within the lever 218 defined by the slot 222 provided in the lever 218. The transverse member 220 is mounted on a spring (not shown), located between the transverse member 220 and the end region of the lever 218 opposed to that end region of the lever 218 mounted on the pivot 219.

In use, a sample support (microscope slide, 6) is positioned in a machined recess (not shown) provided on the sample support holding member 202, the sample bearing surface facing outwards, away from the support holding member 202. The angled arms 206 (and thus, the head-piece 210, and the support retaining member 216) are moved into the closed position by the operator, as shown in Figure 15b. In the closed position a support cell is formed, comprising a substantially sealed chamber defined by the sample bearing surface of the sample support 6, the support retaining member 216, and the O ring gasket 224 provided in a machined recess formed in the surface of the support retaining member 216. A good seal between the gasket 224 and the sample-bearing surface of the sample support 6 is ensured by the spring-loaded mounting pins 212, 214, which bias the support retaining member 216 towards the support holding member 202 and the support 6.

Once in the closed position, the arrangement is clamped by the clamping lever 218. To effect clamping the lever 218 is moved upwards about the pivot 203. The transverse member 220 engages a co-operating recess 226 in the angled arms 206, such that the upward movement of the clamping lever 218 forces the transverse member to move within the slot 222 away from the pivot 219, against the spring located within the lever 218, causing the spring to become compressed. The compression of the spring provides the clamping force which clamps the arrangement in the closed position.

When the sample has been processed, the clamping lever 218 may be moved downwards about the pivot 219, disengaging the transverse member 220 from the co-operating recess 226. The support retaining member 216 may then be moved away from the support holding member 202, and the support 6 removed from the apparatus. If further samples are to be processed, the procedure may then be repeated.

Figure 16 illustrates an array of six support cells of the type shown in Figure 15. The



support cells are arranged in three pairs, each pair being associated with a respective support holding member 202, such that one thermoregulatable plate comprised in the support holding member 202 may regulate the temperature of two sample supports 6. Each support cell is provided with a fluid inlet 228 and a fluid outlet 230 formed in the support retaining members 216.

The support retaining members 216 are provided with round-bottomed grooves where the inlet 228 and outlet 230 enter the support cell, which grooves allow for even distribution of fluid across the width of the sample support as fluid enters the support cell, and which facilitate egress of the fluid via outlet 230. The apparatus comprises fluid delivery means, including ten reservoirs of fluids accommodated within the housing 200. Capillary tubing conducts fluids from the reservoirs, through apertures in the housing and through bores in the support retaining members 216 to the support cell inlets 228. Waste fluid egresses from the outlets 230, through bores in the support retaining members 216 and thence via capillary tubing through apertures in the housing 200 to a waste fluid collecting vessel. Conveniently the capillary tubing is attached to, or located within, the angled arms 206, making for a tidier arrangement with less chance of the tubing becoming snagged.

In a preferred embodiment, the automated apparatus for processing samples comprises four arrays of support holding members of the type shown in Figure 16, such that the apparatus may process up to 24 samples simultaneously. The arrangement illustrated in Figure 16 has the advantage that an operator may observe the processing fluids within the support cells (the support retaining members 216 comprising perspex) and thus easily monitor the progress of the sample processing.

The preferred embodiment of the automated apparatus is provided with fluid delivery means of the type illustrated schematically in Figure 17. The fluid delivery means comprises two Kloen syringe pumps 240a, b fitted with 1ml syringes. At the top of each syringe is located a 6-port rotary valve 242a, b. each rotary valve 242a, b being provided with 5 inlet ports and 1 exit port. Thus syringe pump 240a may pump fluid selected from one of five different fluid reservoirs 1-5, and syringe pump 240b may pump fluid selected from one of five different fluid reservoirs 6-10, each reservoir being connected to the



syringe pumps 240a, b by a respective port in the rotary valves 242a, b. The syringe pumps 240a, b force fluid via the exit ports of the rotary valves 242a, b into a common mixing valve 244, thence to a distribution valve 246, which directs the fluid to the appropriate array of support cells. Each support cell is provided with a 3-way valve, such that respective support cells may be processed individually with different processing fluids if desired, and allowing for the manual introduction of processing fluids into individual support cells (e.g. by means of a micropipette).

Tables I and II describe example protocols for performing haematoxylin/eosin staining and in situ hybridisation respectively using automated apparatus according to the invention.

It will be clear that other embodiments of the invention could be made and such variants are intended to fall within the scope of the present invention.



TABLE I

**Haematoxylin and Eosin Stain**  
*Example Protocol for Slide Processor*

Stage	Pump No.	Soln.	Ratio	Vol. ( $\mu$ l)	Rate (ml/min)	Cells	Pause (sec)	Repeat	Temp. (°C)
1	1	EtOH	-	1000	2	1-6	60	2	22
2	1 & 2	EtOH/H <sub>2</sub> O	9:1	1000	2	1-6	60	1	22
3	1 & 2	EtOH/H <sub>2</sub> O	7:3	1000	2	1-6	60	1	22
4	1 & 2	EtOH/H <sub>2</sub> O	1:1	1000	2	1-6	60	1	22
5	2	H <sub>2</sub> O	-	1000	2	1-6	-	10	22
6	3	Haem.	-	800	2	1-6	300	1	22
7	2	H <sub>2</sub> O	-	1000	2	1-6	10	10	22
8	4	H+EtOH	-	1000	2	1-6	30	3	22
9	2	H <sub>2</sub> O	-	1000	2	1-6	-	10	22
10	5	Eosin	-	1000	2	1-6	120	1	22
11	2	H <sub>2</sub> O	-	1000	2	1-6	-	15	22
12	1 & 2	EtOH/H <sub>2</sub> O	1:1	1000	2	1-6	60	1	22
13	1 & 2	EtOH/H <sub>2</sub> O	7:3	1000	2	1-6	60	1	22
14	1 & 2	EtOH/H <sub>2</sub> O	9:1	1000	2	1-6	60	2	22
15	1	EtOH	-	1000	2	1-6	60	4	22
16	6	Xylene	-	100	2	1-6	30	4	22



TABLE II

**In Situ Hybridisation from Paraffin Section**  
*Example Protocol for Slide Processor*

Stage	Pump No.	Soln.	Ratio	Vol. ( $\mu$ l)	Rate (ml/min)	Cells	Pause	Repeat	Temp. (°C)
1	1	Xylene	-	1000	2	1-8	00:60	4	37
2	1	EtOH	-	1000	2	1-8	00:60	2	22
3	2 & 3	EtOH/H <sub>2</sub> O	9:1	1000	2	1-8	00:60	1	22
4	2 & 3	EtOH/H <sub>2</sub> O	7:3	1000	2	1-8	00:60	1	22
5	2 & 3	EtOH/H <sub>2</sub> O	1:1	1000	2	1-8	00:60	1	22
6	3	H <sub>2</sub> O	-	1000	2	1-8	-	10	22
7	4	Preflyb. Buffer	-	300	2	1-8	60:00	1	42
8	Hold	Manual add. probe					360:00		42
9	5	1 x SSC	-	1000	2	1-8	30	5	22
10	5	1 x SSC	-	1000	2	1-8	30	20	55
11	6	0.1 x SSC	-	1000	2	1-8	-	2	22
12	2 & 3	EtOH/H <sub>2</sub> O	1:1	1000	2	1-8	60	2	22
13	2 & 3	EtOH/H <sub>2</sub> O	7:3	1000	2	1-8	60	2	22
14	2 & 3	EtOH/H <sub>2</sub> O	9:1	1000	2	1-8	60	2	22
15	2	EtOH	-	1000	2	1-8	60	2	22



## CLAIMS

1. A support retaining member for use in processing a sample supported on a support, the member being such that when assembled together with a support it forms a support cell, the support cell comprising a substantially sealed chamber, the chamber being provided with a fluid inlet and a fluid outlet for the introduction and removal respectively of fluids used in processing the sample.
2. A support retaining member according to claim 1, wherein the support is a microscope slide.
3. A support retaining member according to claim 1 or 2, wherein part of the internal surface of the substantially sealed chamber is defined by the sample bearing surface of the support.
4. A support retaining member according to any one of claims 1, 2 or 3, comprising a synthetic moulded plastics material.
5. A support retaining member according to any one of the preceding claims, comprising sealing means to assist in the formation of a substantially sealed chamber.
6. A support retaining member according to any one of the preceding claims, comprising a gasket of silicon rubber or similar material.
7. A support retaining member according to any one of the preceding claims, adapted for use with clamping means, which clamping means serves to clamp together the components of the support cell.
8. A support retaining member according to claim 7, wherein the clamping means comprises biasing means.
9. A support retaining member according to any one of the preceding claims, comprising



spring biasing means.

10. A support retaining member according to any one of the preceding claims, adapted for use in a support cell comprising temperature regulation means to regulate the temperature of the sample.

11. A support retaining member according to any one of the preceding claims, adapted for use in a support cell for performing the polymerase chain reaction (PCR) on a sample.

12. A support retaining member according to any one of the preceding claims, adapted for use in a support cell having a substantially sealed chamber volume in the range 50-300 $\mu$ l.

13. A support retaining member according to any one of the preceding claims, in combination with a support holding member.

14. A support cell comprising a support retaining member according to any one of claims 1-12.

15. Automated apparatus for processing a sample supported on a support, the apparatus comprising: support holding means for holding one or more supports, the sample on the or each support being present within a respective substantially sealed chamber; fluid delivery for delivery processing fluid to the or each chamber; waste fluid collecting means for removing fluid from the or each chamber; and computer control means.

16. Apparatus according to claim 15, for use with a support cell according to claim 14.

17. Apparatus according to claim 15 or 16, comprising clamping means for clamping together the components of the support cell.

18. Apparatus according to any one of claims 15, 16 or 17 wherein the fluid delivery



means comprises one or more syringe pumps.

19. Apparatus according to any one of claims 16, 17 or 18, wherein each support cell is provided with an individual inlet valve close to a fluid inlet provided in the support cell.

20. Apparatus according to any one of claims 15-19, comprising temperature control means.

21. Apparatus according to any one of claims 15-20, comprising a Peltier device.

22. A method of processing a sample on a support, comprising use of a support cell according to claim 14.

23. A method of processing a sample on a support, comprising use of automated apparatus according to any one of claims 15-21.

24. A method of making a support cell, comprising assembling a support together with a support retaining member according to any one of claims 1-12.

25. A method according to claim 24, wherein the components of the support cell are held together by clamping means and/or biasing means.



1/16

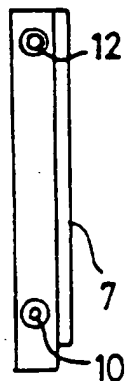


Fig. 1b

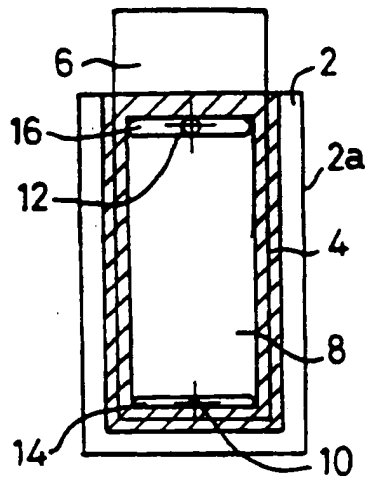


Fig. 1a

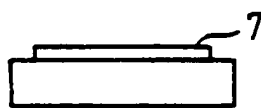


Fig. 1c

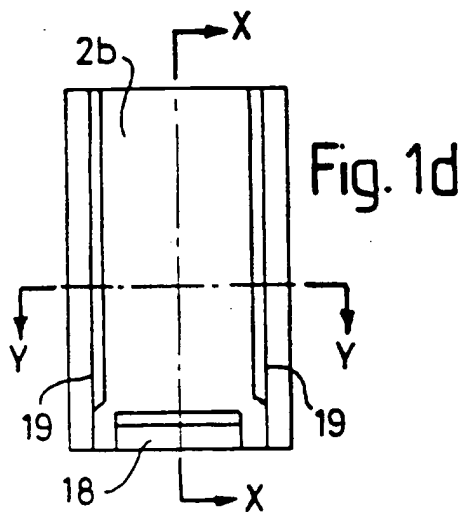


Fig. 1d

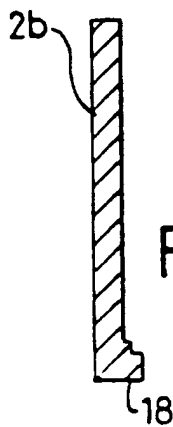


Fig. 1f

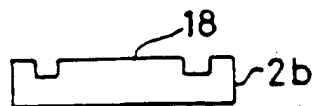


Fig. 1e



Fig. 1g

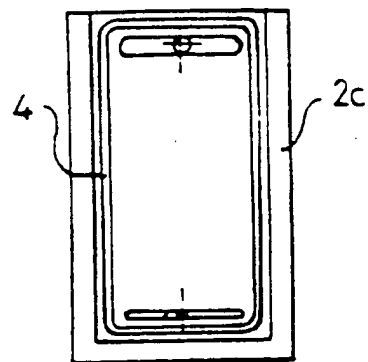


Fig. 2



2/16

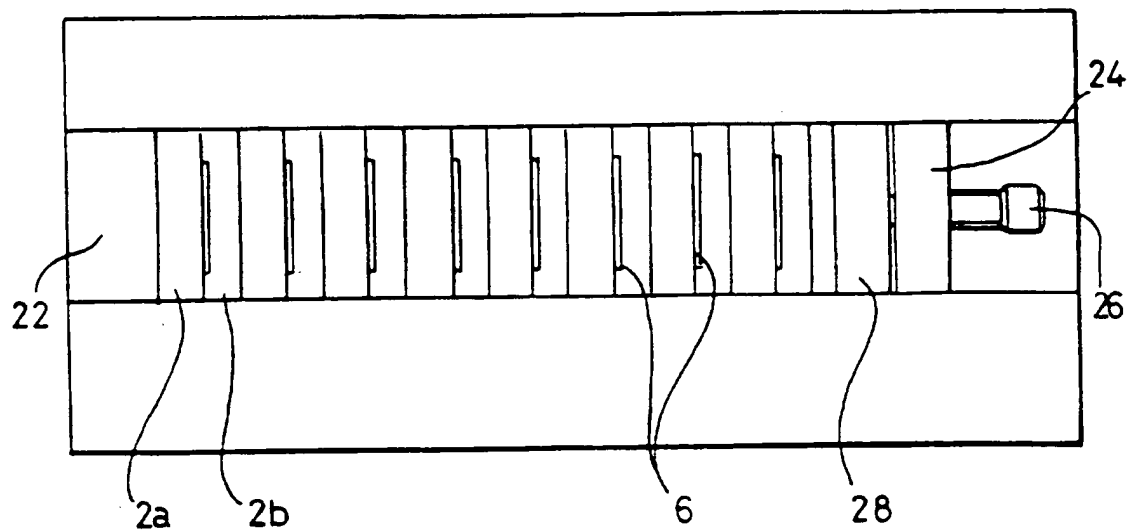
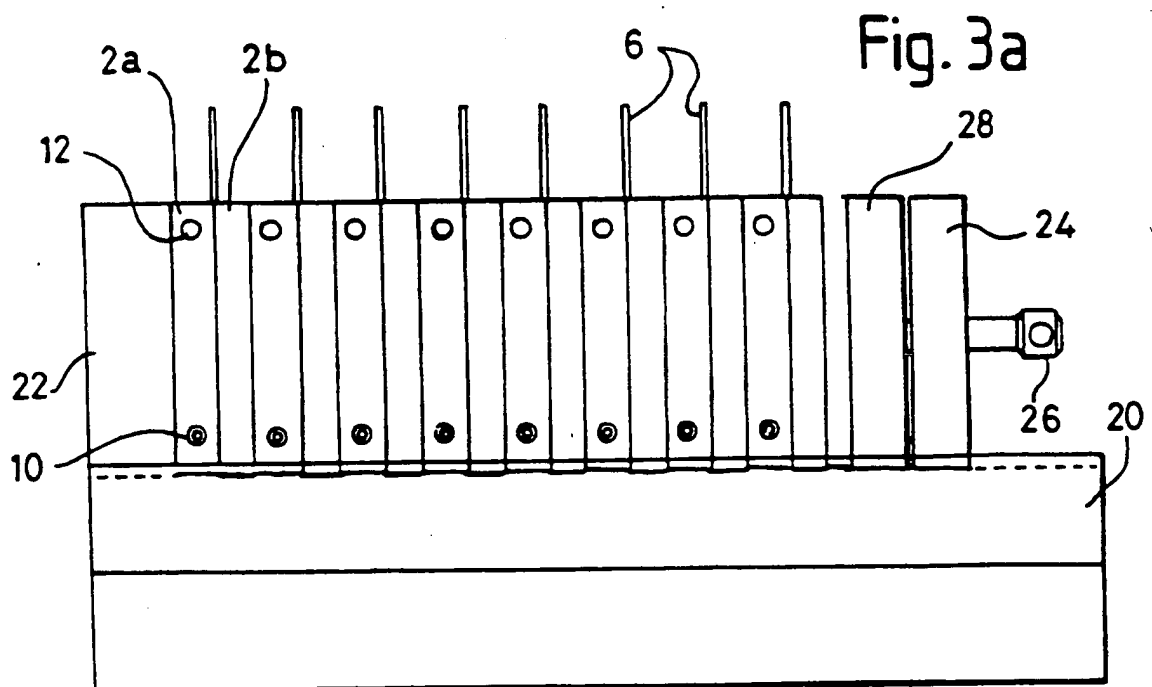


Fig. 3b



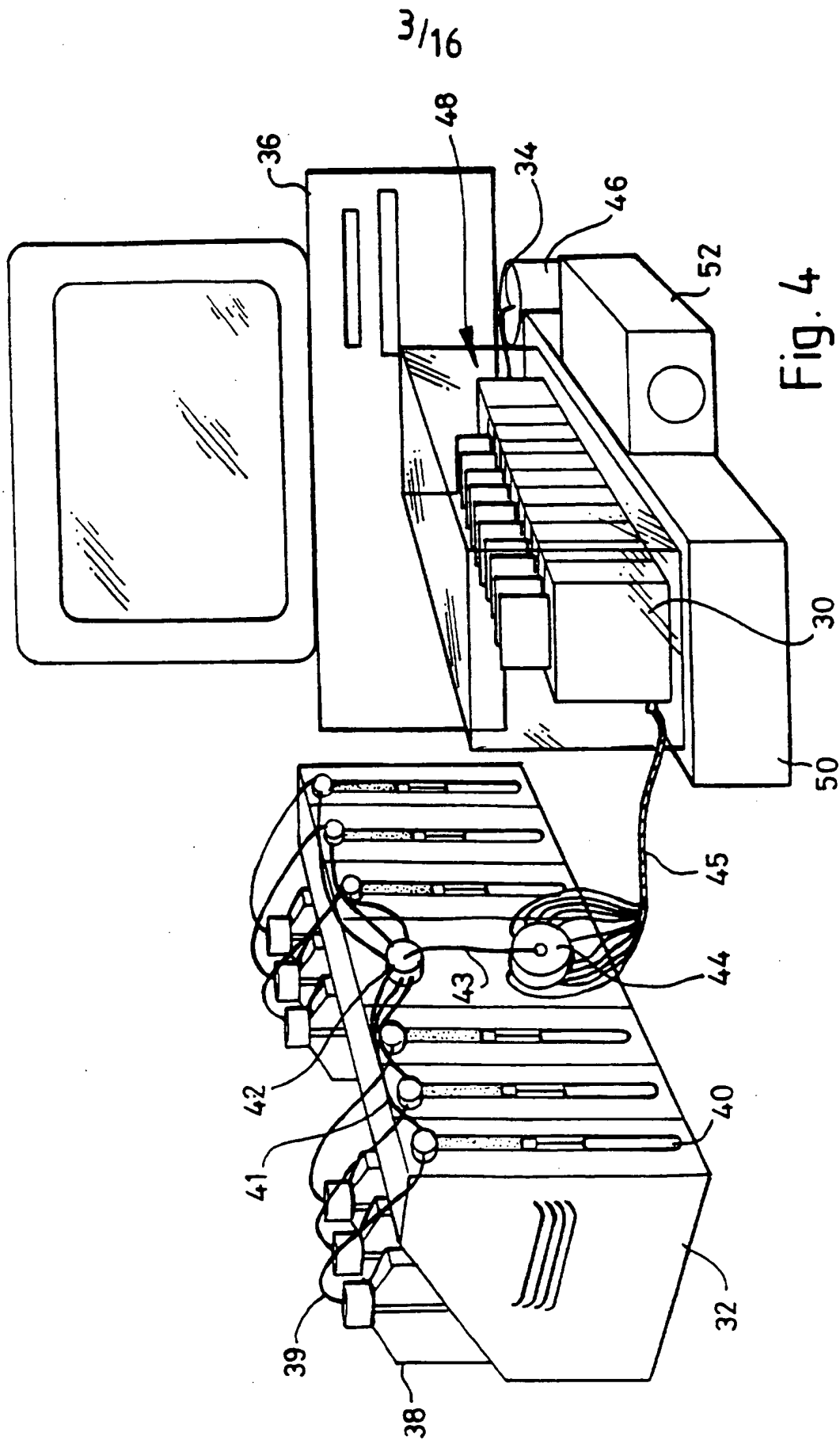
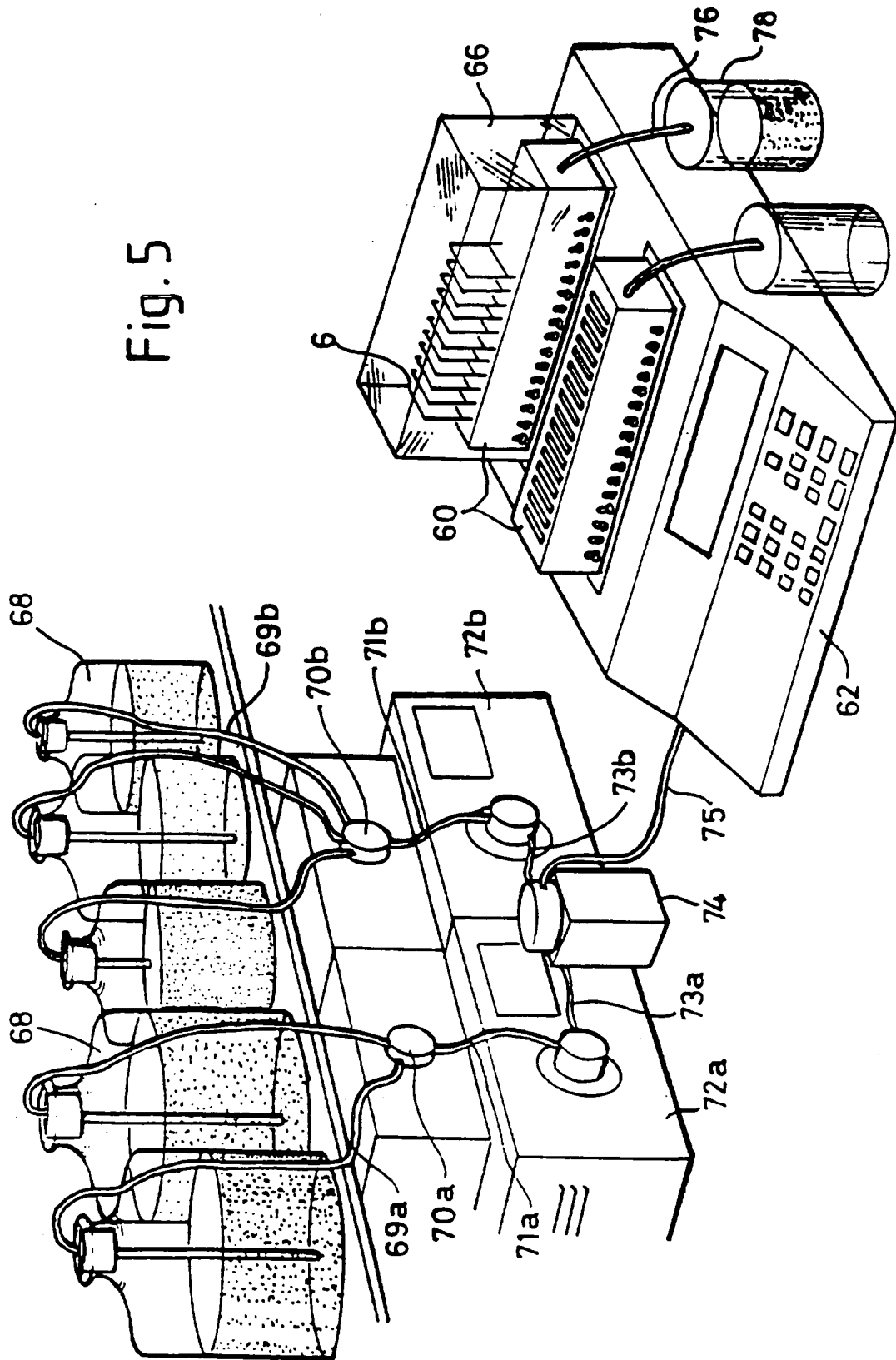


Fig. 4



4/16

Fig. 5





5/16

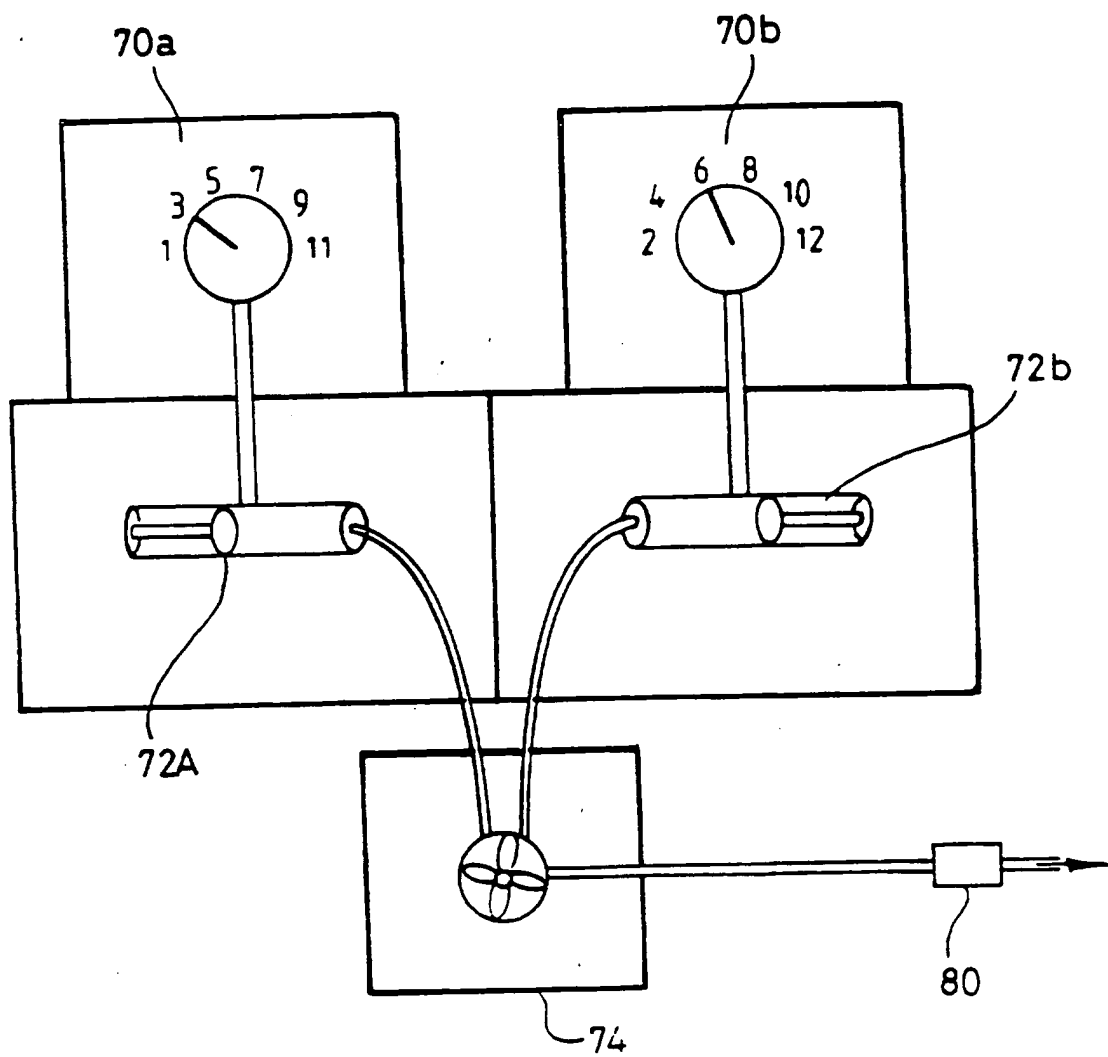


Fig. 6



6/16

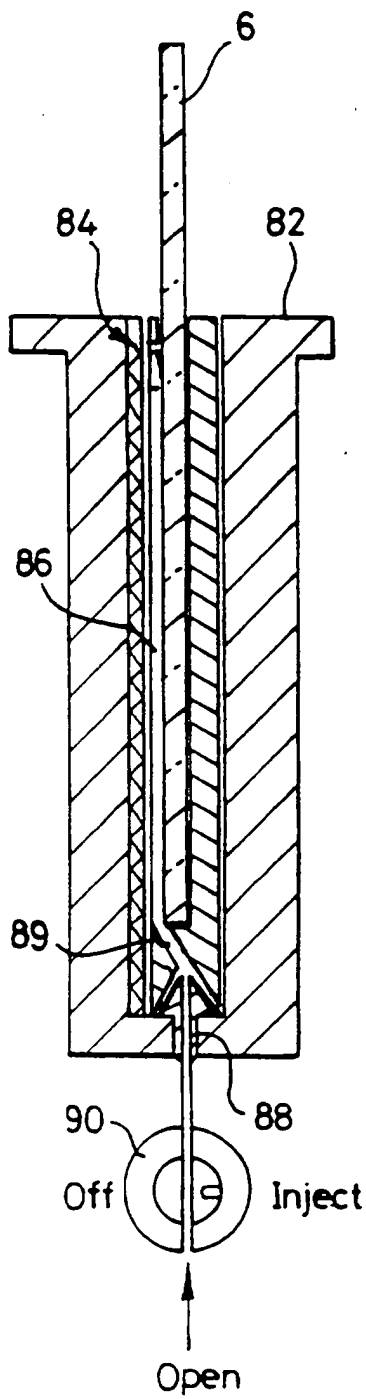


Fig. 7a

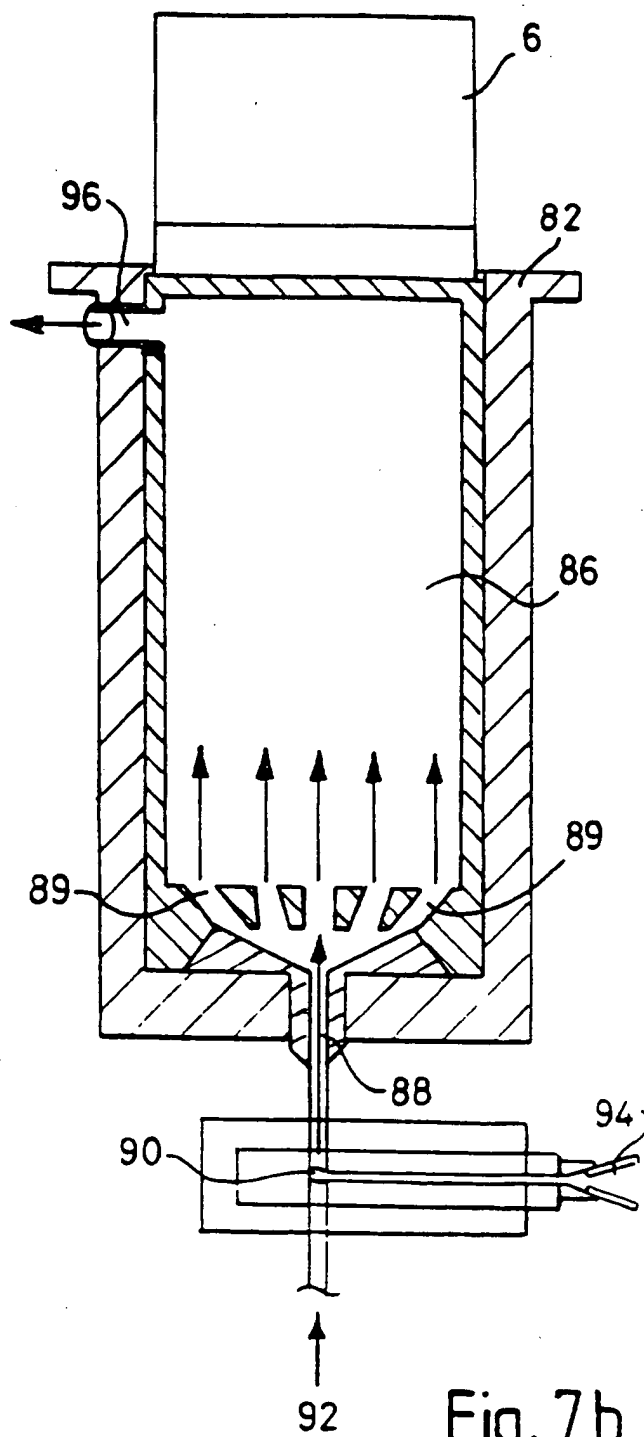


Fig. 7b



# Flow Diagrams of Histology

A → B → C → D → E → F → G → H → I → J → K → L

## Haemotoxylin and Eosin

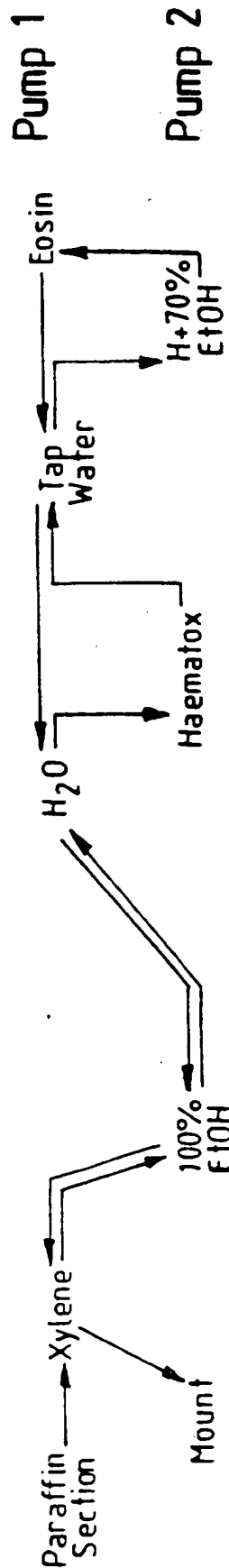
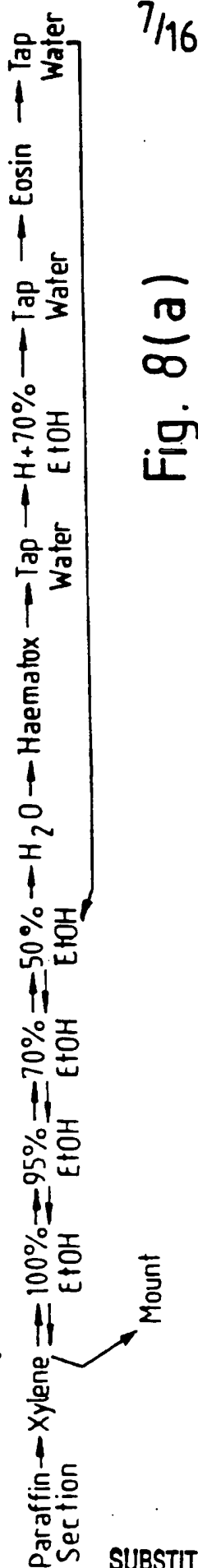


Fig. 8(b)



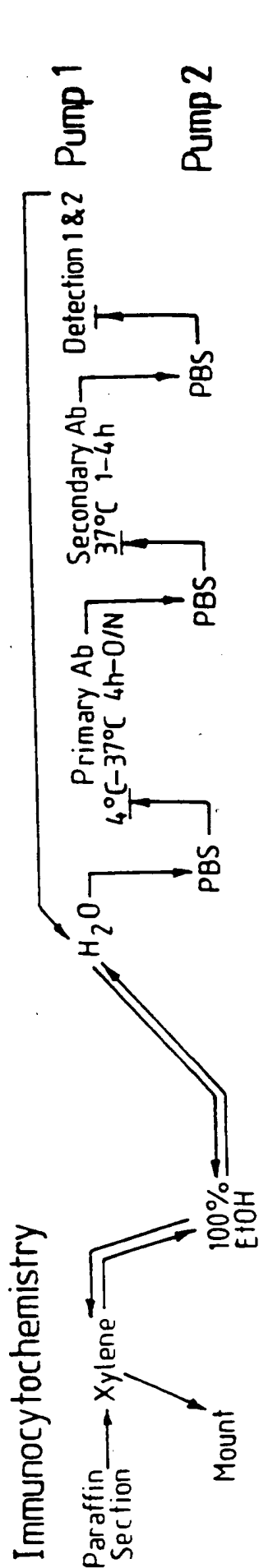


Fig. 9 (a)

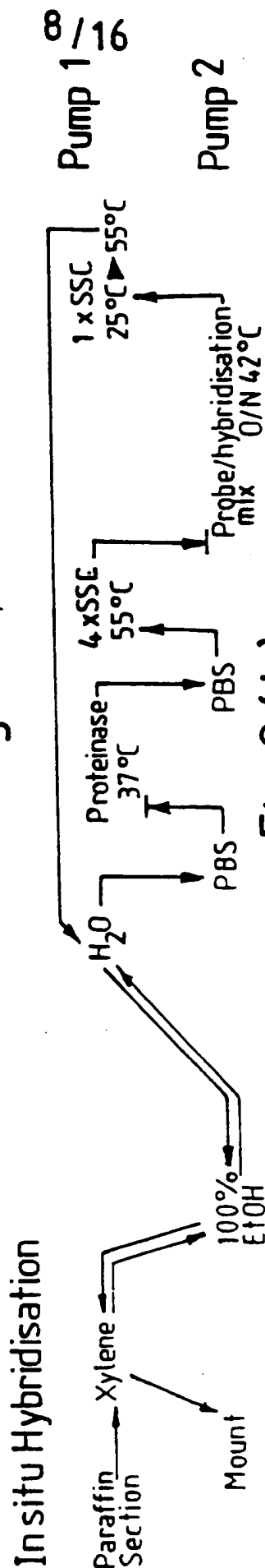


Fig. 9 (b)

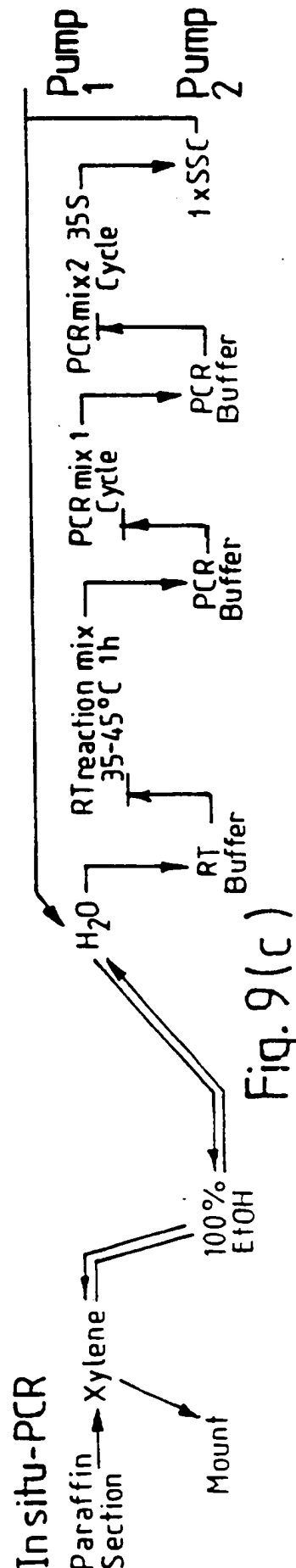


Fig. 9 (c)



9/16

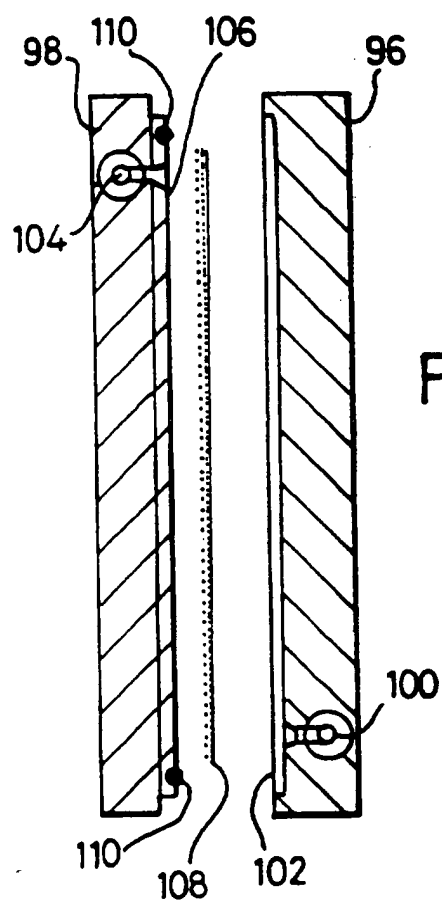


Fig. 10a

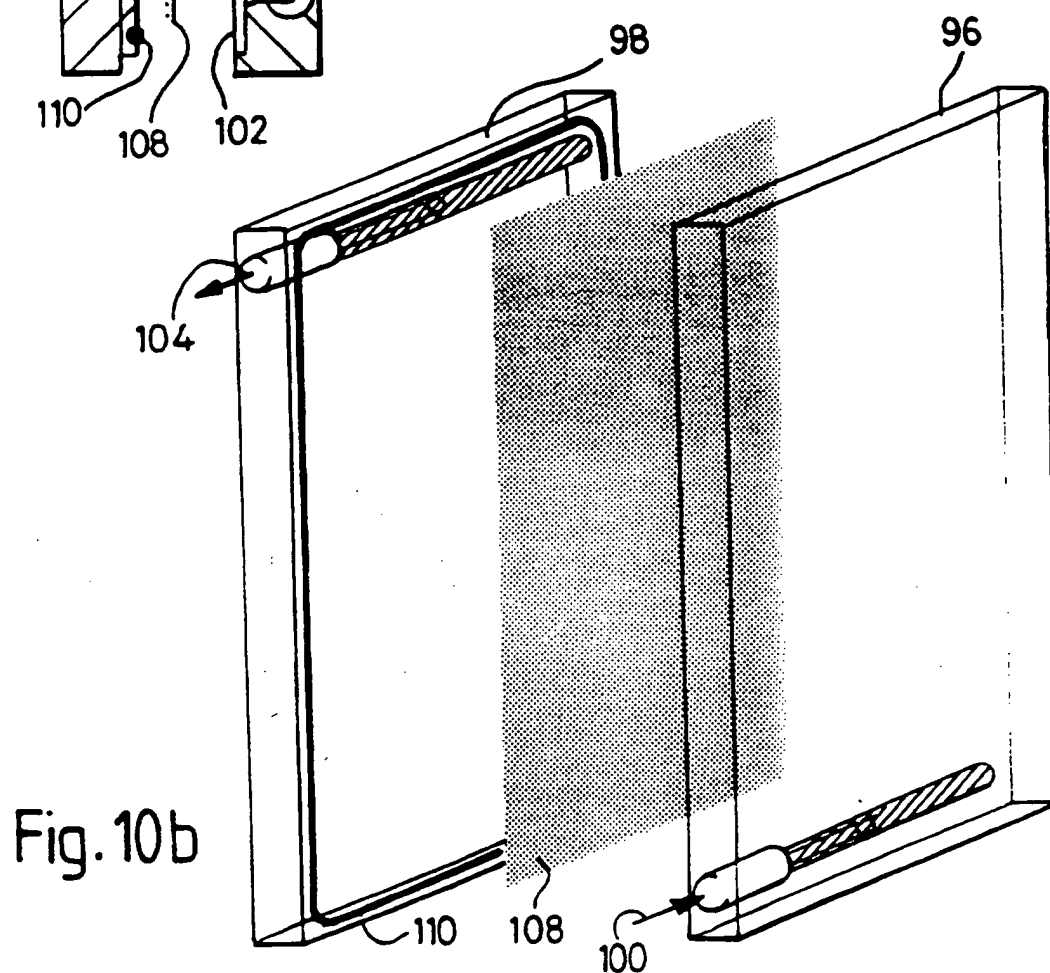


Fig. 10b



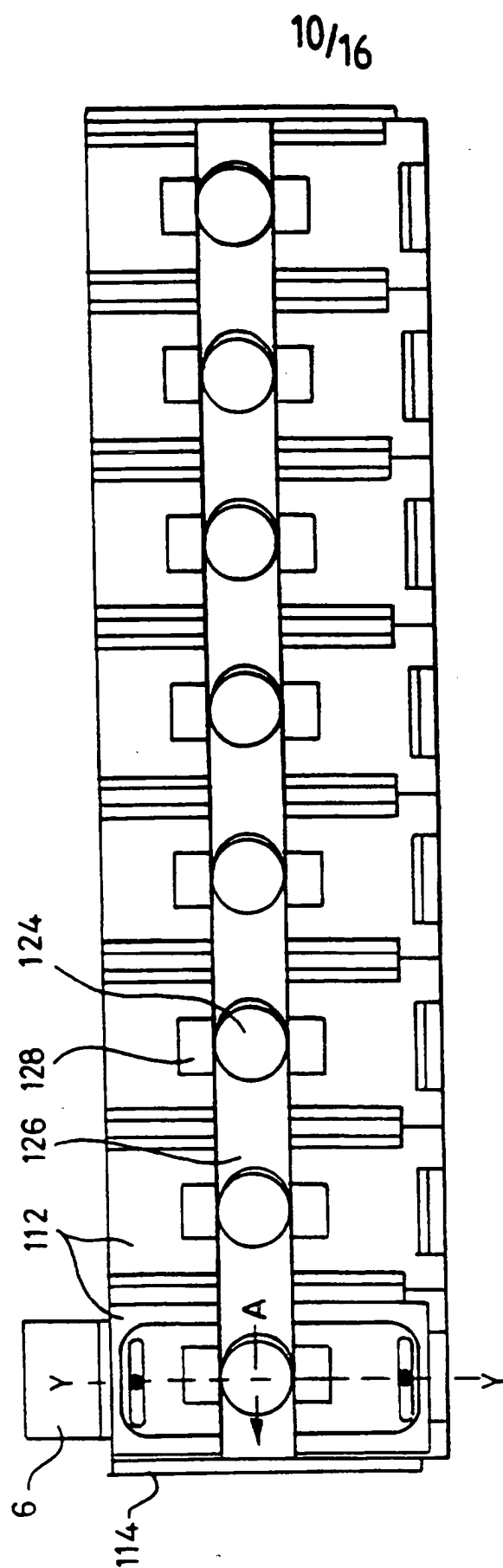


Fig. 11a



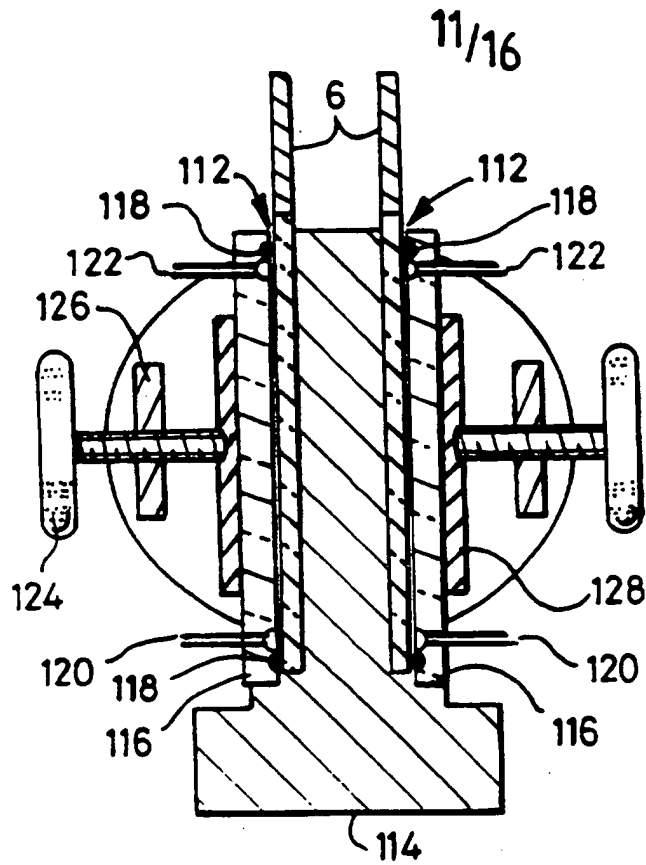


Fig. 11b

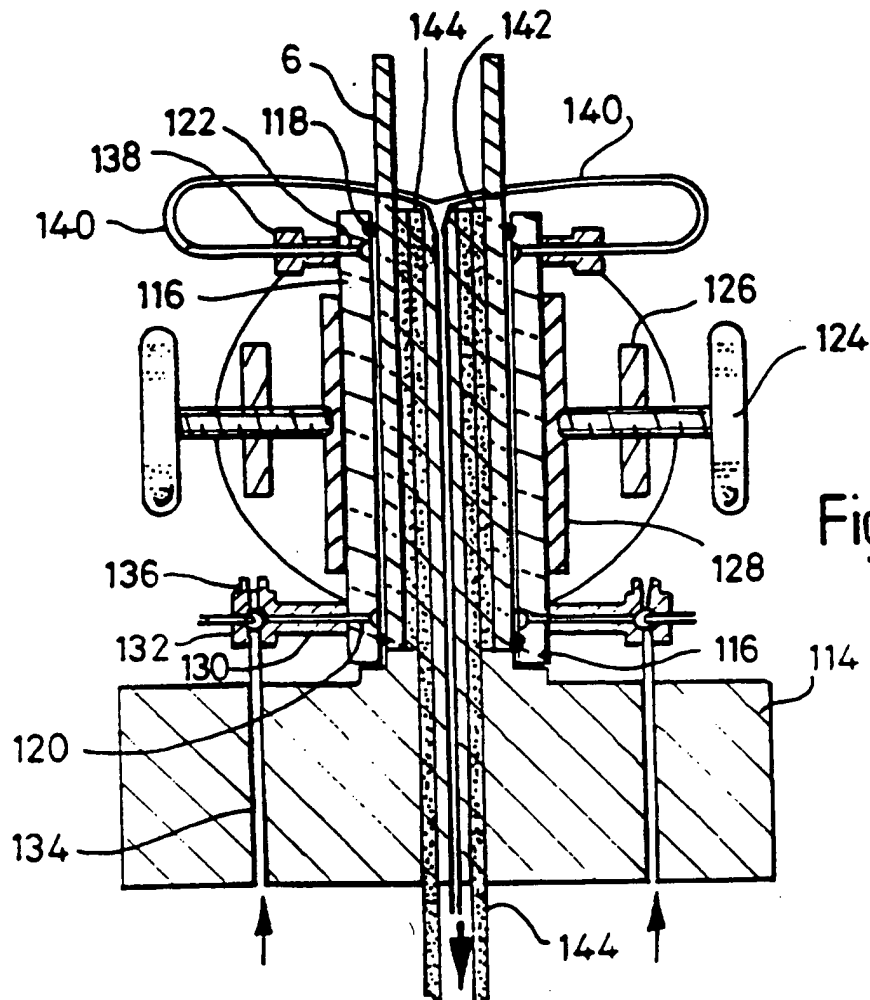


Fig. 12



12/16

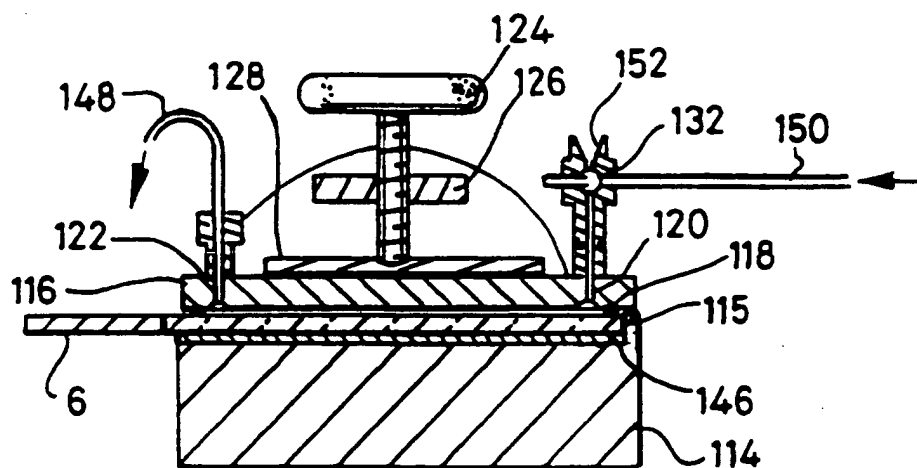


Fig. 13a

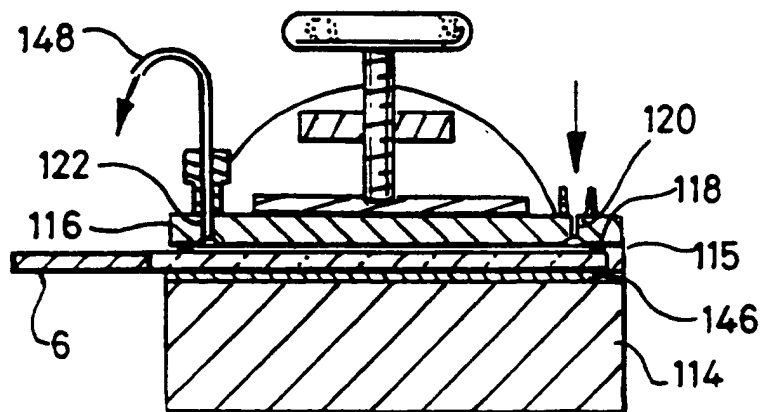


Fig. 13b



13/16

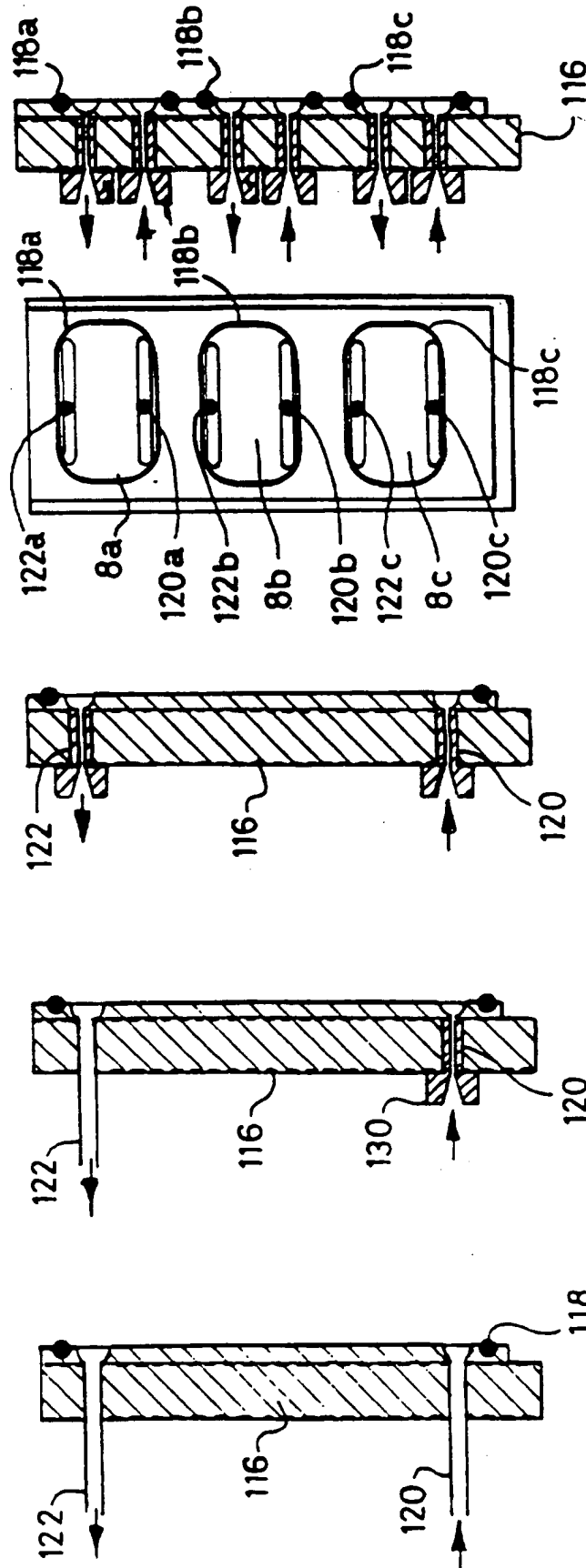
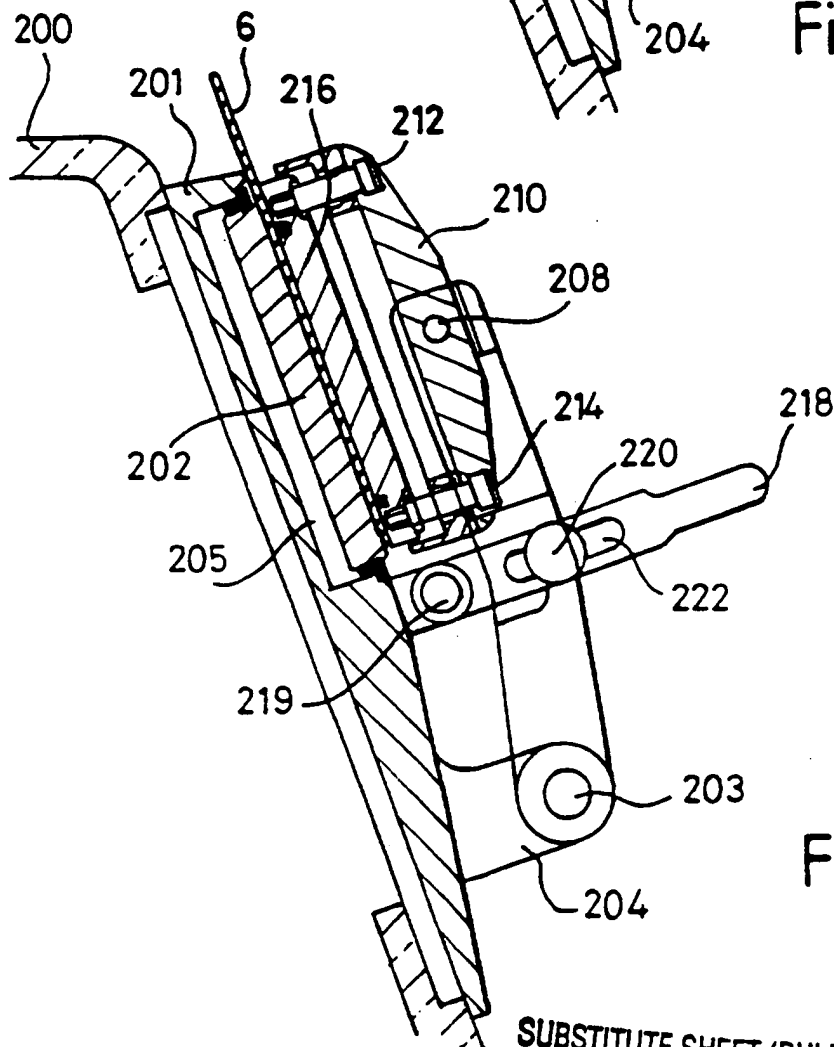
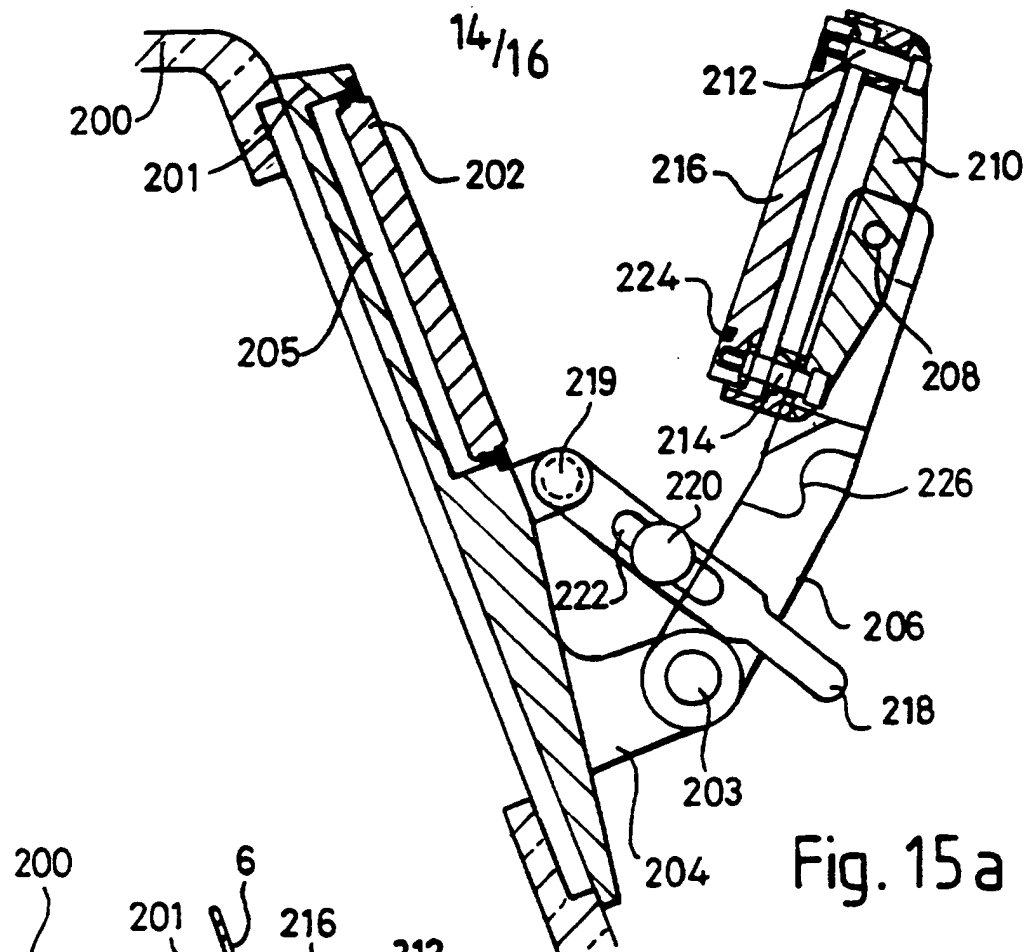


Fig. 14(a) Fig. 14(b) Fig. 14(c) Fig. 14(d) Fig. 14(e)







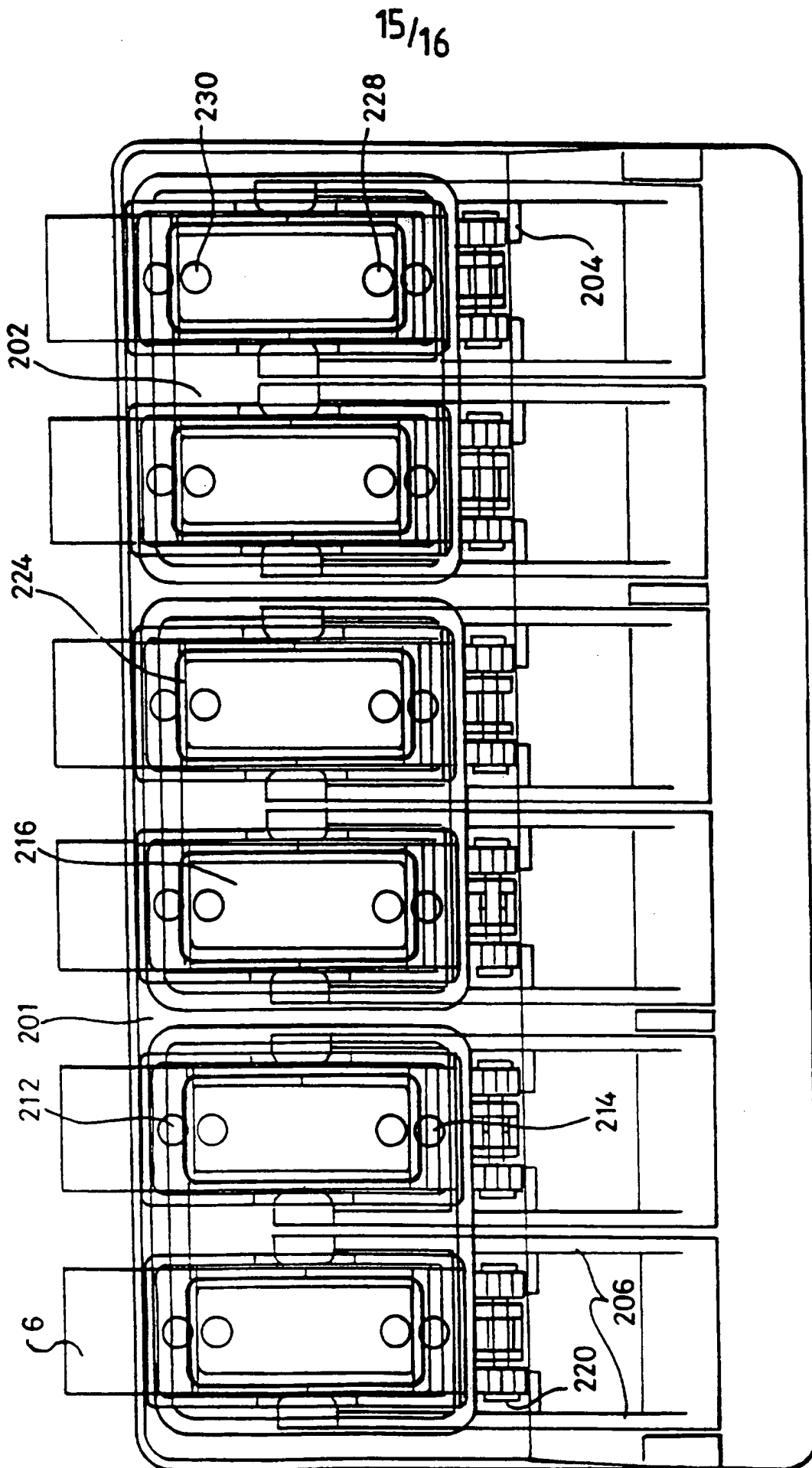


Fig. 16



16/16

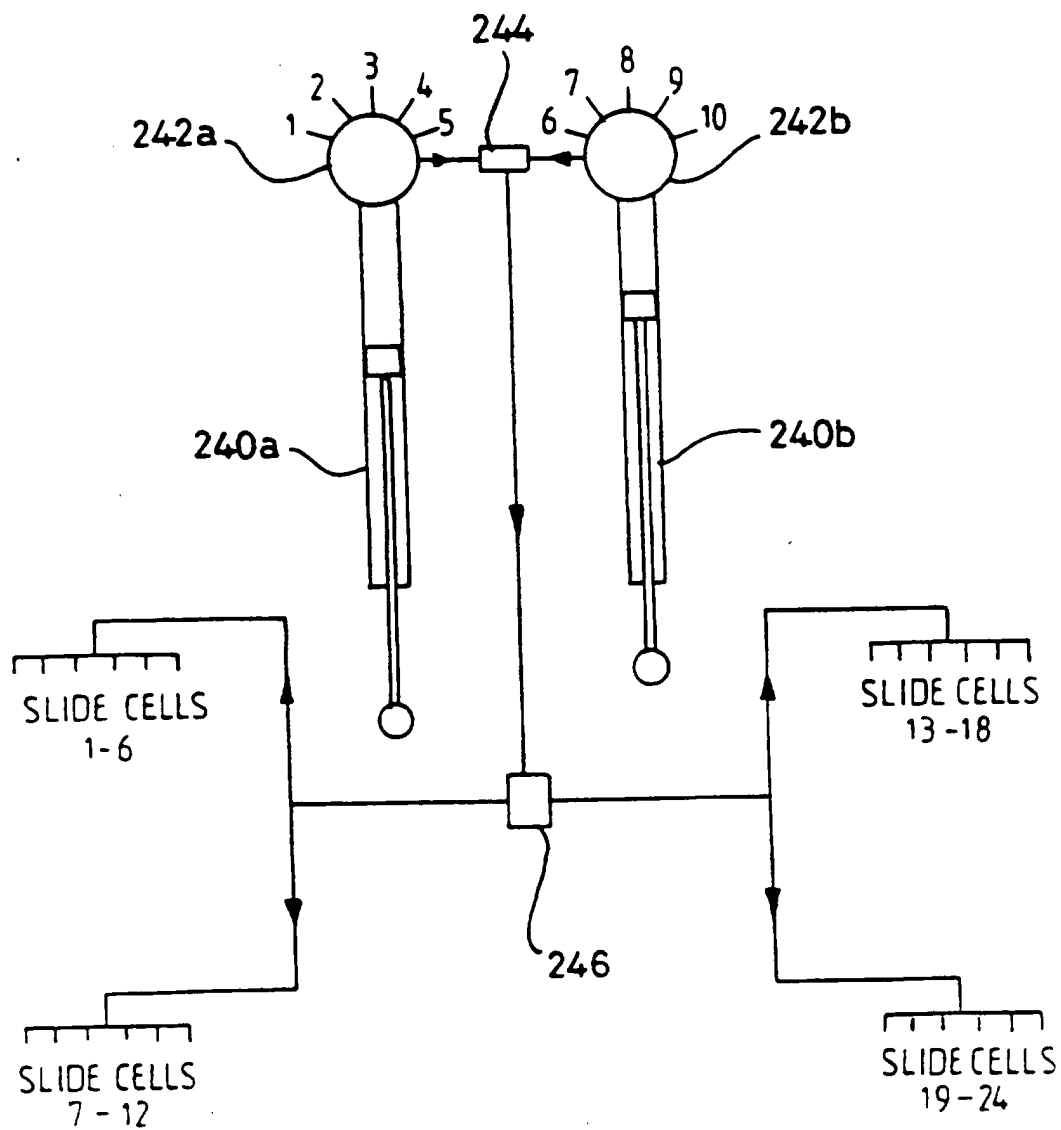


Fig. 17



# INTERNATIONAL SEARCH REPORT

Int. Appl. No.  
PCT/GB 96/00758

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 6 B01L7/00 G01N1/31 C12Q1/68

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 G01N B01L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO,A,89 00887 (BOGEN STEVEN ALLEN) 9 February 1989  see page 10, line 10 - line 16 see page 12, line 31 - page 13, line 11; figures see page 18, line 29 - line 32 ---	1-9, 13-19, 22-25
X	US,A,5 281 516 (STAPLETON MARILYN J ET AL) 25 January 1994 see column 7, line 38 - column 8, line 4 see column 9, line 7 - line 15; figure 5 see column 13, line 66 - column 14, line 48 ---	1-3,10, 11,20
X	---	21
	-/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

### \* Special categories of cited documents:

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

- \*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- \*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- \*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- \*A\* document member of the same patent family

Date of the actual completion of the international search

5 July 1996

Date of mailing of the international search report

12. 07. 96

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl,  
Fax (+ 31-70) 340-3016

Authorized officer

Hocquet, A



## INTERNATIONAL SEARCH REPORT

Inter .al Application No  
PCT/GB 96/00758

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	GB,A,2 261 111 (STUART SCIENTIFIC COMPANY) 5 May 1993 see page 6, line 20 - page 7, line 32; figure 1 ---	21
X	EP,A,0 464 861 (UNIVERSITY OF LEICESTER) 8 January 1992 see column 5, line 14 - line 53; figures ---	1
A	US,A,5 192 503 (MCGRATH CHARLES M ET AL) 9 March 1993 see column 1, line 40 - line 45 see column 6, line 62 - column 7, line 24; figures 6,8 see column 11, line 45 - column 12, line 9; figure 9 see column 12, line 51 - line 54 ---	1-25 12
A	EP,A,0 310 399 (SHANDON SCIENT LTD) 5 April 1989 see column 3, line 43 - column 4, line 9; figures 10-12,15 ---	13,15
A	WO,A,91 07486 (STAPLETON) 30 May 1991 see page 18, line 22 - page 19, line 37; figures 5,12 see page 23, line 17 - page 24, line 10; figures 10,11 ---	1
A	WO,A,93 19207 (GENE TEC) 30 September 1993 see page 18, line 9 - line 29; figure 11 -----	1



# INTERNATIONAL SEARCH REPORT

Inte mal Application No  
PCT/GB 96/00758

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO-A-8900887	09-02-89	US-A- 4847208 US-A- 5073504	11-07-89 17-12-91
US-A-5281516	25-01-94	US-A- 5188963 EP-A- 0632839 US-A- 5346672 WO-A- 9319207 US-A- 5382511 AU-B- 7787291 CA-A- 2068891 EP-A- 0502108 JP-T- 5501647 WO-A- 9107486 US-A- 5436129 US-A- 5451500	23-02-93 11-01-95 13-09-94 30-09-93 17-01-95 13-06-91 18-05-91 09-09-92 02-04-93 30-05-91 25-07-95 19-09-95
GB-A-2261111	05-05-93	NONE	
EP-A-464861	08-01-92	EP-A- 0270363 GB-A- 2200210 GB-A- 2234348 JP-A- 63214648	08-06-88 27-07-88 30-01-91 07-09-88
US-A-5192503	09-03-93	NONE	
EP-A-0310399	05-04-89	JP-A- 1257261 US-A- 4985206	13-10-89 15-01-91
WO-A-9107486	30-05-91	US-A- 5188963 AU-B- 7787291 CA-A- 2068891 EP-A- 0502108 JP-T- 5501647 US-A- 5346672 US-A- 5436129 US-A- 5451500 US-A- 5281516	23-02-93 13-06-91 18-05-91 09-09-92 02-04-93 13-09-94 25-07-95 19-09-95 25-01-94
WO-A-9319207	30-09-93	US-A- 5346672 US-A- 5281516	13-09-94 25-01-94



# INTERNATIONAL SEARCH REPORT

Internal Application No

PCT/GB 96/00758

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
W0-A-9319207		EP-A- 0632839	11-01-95
-----			



**This Page is Inserted by IFW Indexing and Scanning  
Operations and is not part of the Official Record**

**BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ BLACK BORDERS
- ☐ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES
- ☒ FADED TEXT OR DRAWING
- ☐ BLURRED OR ILLEGIBLE TEXT OR DRAWING
- ☐ SKEWED/SLANTED IMAGES
- ☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS
- ☐ GRAY SCALE DOCUMENTS
- ☒ LINES OR MARKS ON ORIGINAL DOCUMENT
- ☐ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY
- ☐ OTHER: \_\_\_\_\_

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.**